

OM of: US-09-439-311-2 to: Issued\_Patents\_NA:\* out\_format : pfs  
 Date: Apr 17, 2002 3:10 AM

About: Results were produced by the GenCore software, version 4.5,  
 Copyright (c) 1993-2000 Compugen Ltd.

## Command line parameters:

-MODE=frame+27n model -DEV=xlp  
 -O=/cg2\_6/prodata/2/lna/5A\_COMB.seq:US-09-358-972-102  
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 -GAPEXT=4.000 -MINMATCH=0.100 -LOOPEXT=0.000  
 -GAPOP=4.500 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500  
 -DELOP=6.000 -EGAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500  
 -TRANS=human40.cdi -LIST=45 -DOCALLIGN=200 -THR\_SCORE=pcr  
 -THR\_MAX=100 -THR\_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pfs  
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## Search information block:

Query: US-09-439-311-2  
 Query length: 333  
 Database: Issued\_Patents\_NA\*  
 Database sequences: 351203  
 Database length: 11323899  
 Search time (sec): 85.700000

## Score\_list:

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APPLICANT: Welch, Roy
TITLE OF INVENTION: Nucleic Acid Detection
FILE REFERENCE: Pro-103 6868/75528
CURRENT APPLICATION NUMBER: US/09/358,972
CURRENT FILING DATE: 1999-07-22
EARLIER APPLICATION NUMBER: 09/252,436
EARLIER FILING DATE: 1999-02-18
EARLIER APPLICATION NUMBER: 09/042,287
EARLIER FILING DATE: 1998-03-13
NUMBER OF SEQ ID NOS: 290
SOFTWARE: Patentln Ver. 2.0
SEQ ID NO 103
LENGTH: 30
TYPE: DNA
ORGANISM: Campylobacter jejuni
FEATURE:
OTHER INFORMATION: probe to Campylobacter jejuni
US-09-358-972-103
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; Patent No. 6270974
; GENERAL INFORMATION:
; APPLICANT: Shultz, John W
; APPLICANT: Lewis, Martin K
; APPLICANT: Leipzig, Donna
; APPLICANT: Mandrekas, Michelle
; APPLICANT: Kephart, Daniel
; APPLICANT: Rhodes, Richard B
; APPLICANT: Andrews, Christine A
; APPLICANT: Hartnett, James R
; APPLICANT: Gu, Trent
; APPLICANT: Wood, Keith V
; TITLE OF INVENTION: EXOGENOUS NUCLEIC ACID DETECTION
; FILE REFERENCE: EXOGENOUS NUCLEIC ACID DETECTION
; CURRENT APPLICATION NUMBER: US/09/406,147
; CURRENT FILING DATE: 1999-09-27
; EARLIER APPLICATION NUMBER: 09/252,436
; EARLIER FILING DATE: 1999-02-18
; EARLIER APPLICATION NUMBER: 09/042,287
; EARLIER FILING DATE: 1998-03-13
; NUMBER OF SEQ ID NOS: 92
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 32
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; TYPE: DNA
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; Patent No. 6270974
; GENERAL INFORMATION:
; APPLICANT: Shultz, John W
; APPLICANT: Lewis, Martin K
; APPLICANT: Leipzig, Donna
; APPLICANT: Mandrekas, Michelle
; APPLICANT: Kephart, Daniel
; APPLICANT: Rhodes, Richard B
; APPLICANT: Andrews, Christine A
; APPLICANT: Hartnett, James R
; APPLICANT: Gu, Trent
; APPLICANT: Wood, Keith V
; TITLE OF INVENTION: EXOGENOUS NUCLEIC ACID DETECTION
; FILE REFERENCE: EXOGENOUS NUCLEIC ACID DETECTION
; CURRENT APPLICATION NUMBER: US/09/406,147
; CURRENT FILING DATE: 1999-09-27
; EARLIER APPLICATION NUMBER: 09/252,436
; EARLIER FILING DATE: 1999-02-18
; EARLIER APPLICATION NUMBER: 09/042,287
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; Patent No. 6020163
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; APPLICANT: Conklin, Darrell C.
; TITLE OF INVENTION: LIPOCALIN HOMOLOG
; FILE REFERENCE: 97-24
; CURRENT APPLICATION NUMBER: US/09/130,663A
; CURRENT FILING DATE: 1998-08-05
; EARLIER APPLICATION NUMBER: 60/054,867
; EARLIER FILING DATE: 1997-08-06
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22
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LENGTH: 51  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Oligonucleotide primer: ZC13735.  
US-09-130-663-22

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Quality: 43.00 Length: 16  
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Percent Similarity: 75.000 Percent Identity: 50.000

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||||| :||| |||||:||||| |||:|||||  
1 GGTGAACCTTGACACAGAGATTACAGAGCAGTGTGACACAGGCT 48

seq\_name: /cgn2\_6/protdata/2/lna/6A\_COMB.seq:US-09-081-180-29

## seq\_documentation\_block:

Sequence 29, Application US/09081180

Patent No. 6022847

GENERAL INFORMATION:

APPLICANT: Sheppard, Paul O.

TITLE OF INVENTION: SECRETED SALIVARY ZSIG32

TITLE OF INVENTION: POLYPEPTIDES

NUMBER OF SEQUENCES: 38

CORRESPONDENCE ADDRESS:

ADDRESSEE: Zymogenetics

STREET: 1201 Eastlake Ave. E.

CITY: Seattle

STATE: WA

COUNTRY: USA

ZIP: 98102

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/081,180

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/041,263

FILING DATE: March 19, 1997

ATTORNEY/AGENT INFORMATION:

NAME: Lingenfelter, Susan E

REGISTRATION NUMBER: 41,156

REFERENCE/DOCKET NUMBER: 97-17C1

TELECOMMUNICATION INFORMATION:

TELEPHONE: 206-442-6675

TELEFAX: 206-442-6678

TELEX:

INFORMATION FOR SEQ ID NO: 29:

SEQUENCE CHARACTERISTICS:

LENGTH: 51 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: CDNA

IMMEDIATE SOURCE:

CLONE: ZC13735

US-09-081-180-29

## alignment\_scores:

Quality: 43.00 Length: 16  
Ratio: 3.583 Gaps: 0  
Percent Similarity: 75.000 Percent Identity: 50.000

## alignment\_block:

US-09-439-311-2 x US-09-081-180-29

Align seg 1/1 to: US-09-081-180-29 from: 1 to: 51

253 GlyValValIleGlyLysValAspTyrSerAspGlyAspGluAsnGly 268

||||| :||| |||||:||||| |||:|||||  
1 GGTGAACCTTGACACAGAGATTACAGAGCAGTGTGACACAGGCT 48

seq\_name: /cgn2\_6/protdata/2/lna/6A\_COMB.seq:US-09-040-786-29

## seq\_documentation\_block:

Sequence 29, Application US/09040786

Patent No. 6025197

GENERAL INFORMATION:

APPLICANT: Sheppard, Paul O.

TITLE OF INVENTION: SECRETED SALIVARY ZSIG32

TITLE OF INVENTION: POLYPEPTIDES

NUMBER OF SEQUENCES: 38

CORRESPONDENCE ADDRESS:

ADDRESSEE: Zymogenetics

STREET: 1201 Eastlake Ave. E.

CITY: Seattle

STATE: WA

COUNTRY: USA

ZIP: 98102

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/040,786

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/041,263

FILING DATE: March 19, 1997

ATTORNEY/AGENT INFORMATION:

NAME: Lingenfelter, Susan E

REGISTRATION NUMBER: 41,156

REFERENCE/DOCKET NUMBER: 97-17

TELECOMMUNICATION INFORMATION:

TELEPHONE: 206-442-6675

TELEFAX: 206-442-6678

TELEX:

INFORMATION FOR SEQ ID NO: 29:

SEQUENCE CHARACTERISTICS:

LENGTH: 51 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: CDNA

IMMEDIATE SOURCE:

CLONE: ZC13735

US-09-040-786-29

## alignment\_scores:

Quality: 43.00 Length: 16  
Ratio: 3.583 Gaps: 0  
Percent Similarity: 75.000 Percent Identity: 50.000

## alignment\_block:

US-09-439-311-2 x US-09-040-786-29

Align seg 1/1 to: US-09-040-786-29 from: 1 to: 51

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253 GlyValValIleGlyLysValAspTyrSerAspGlyAspGluAsnGly 268
|||||
1 GGTGAACCTTGACACAGAGATTACAAAGCATGATGACAAAGGCT 48

seq_name: /cgn2_6/ptodata/2/1na/6A_COMB.seq:US-09-432-335-22

seq_documentation_block:
; Sequence 22, Application US/09432335
; Patent No. 6143720
; GENERAL INFORMATION:
; APPLICANT: Conklin, Darrell C.
; TITLE OF INVENTION: LIPOCALIN HOMOLOG
; FILE REFERENCE: 97-24
; CURRENT APPLICATION NUMBER: US/09/432,335
; EARLIER FILING DATE: 1999-11-02
; EARLIER APPLICATION NUMBER: 09/130,663
; EARLIER FILING DATE: 1998-08-06
; EARLIER APPLICATION NUMBER: 60/054,867
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22
; LENGTH: 51
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide primer: ZC13735
US-09-432-335-22

alignment_scores:
Quality: 43.00 Length: 16
Ratio: 3.583 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 50.000

alignment_block:
US-09-439-311-2 x US-09-432-335-22 ..

Align seg 1/1 to: US-09-432-335-22 from: 1 to: 51

253 GlyValValIleGlyLysValAspTyrSerAspGlyAspGluAsnGly 268
|||||
1 GGTGAACCTTGACACAGAGATTACAAAGCATGATGACAAAGGCT 48

seq_name: /cgn2_6/ptodata/2/1na/5A_COMB.seq:US-08-219-012-80

seq_documentation_block:
; Sequence 80, Application US/08219012
; Patent No. 5543293
; GENERAL INFORMATION:
; APPLICANT: Larry Gold
; APPLICANT: Diane Tasset
; TITLE OF INVENTION: Ligands of Thrombin
; NUMBER OF SEQUENCES: 92
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Beaton & Swanson, P.C.
; STREET: 4582 South Ulster Street Parkway, Suite #
; CITY: Denver
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80237
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 5.25 inch, 360 Kb storage
; COMPUTER: IBM compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/219,012
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA: none
; ATTORNEY/AGENT INFORMATION:
```

```
; NAME: Barry J. Swanson
; REGISTRATION NUMBER: 33,215
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (303) 850-9900
; TELEFAX: (303) 850-9401
; INFORMATION FOR SEQ ID NO: 80:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 60 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-219-012-80

alignment_scores:
Quality: 40.00 Length: 19
Ratio: 2.667 Gaps: 0
Percent Similarity: 78.947 Percent Identity: 42.105

alignment_block:
US-09-439-311-2 x US-08-219-012-80 ..

Align seg 1/1 to: US-08-219-012-80 from: 1 to: 60

204 ThrSerValGlyThrGlyLeuGlyAlaLeuAlaGluGluIleAsnArgAs 220
|||||
4 ACCCGGAGCGCGGTGAGGCGTGGAGCGCTGGCCGATGTGTAGCAGCAG 53

220 nAlaAsp 222
|||||
54 CTCGGAT 60

seq_name: /cgn2_6/ptodata/2/1na/6B_COMB.seq:US-08-687-421-268

seq_documentation_block:
; Sequence 268, Application US/08687421
; Patent No. 6177537
; GENERAL INFORMATION:
; APPLICANT: Gold, Larry
; APPLICANT: Janjic, Nebojsa
; APPLICANT: Tasset, Diane
; TITLE OF INVENTION: HIGH-AFFINITY LIGANDS OF BASIC
; TITLE OF INVENTION: FIBROBLAST GROWTH FACTOR AND
; NUMBER OF SEQUENCES: 445
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Swanson & Bratschun, L.L.C.
; STREET: 8400 E. Prentice Avenue, Suite 200
; CITY: Englewood
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB storage
; COMPUTER: IBM compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 6.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/687,421
; FILING DATE: 08-MAY-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/195,005
; FILING DATE: 10-FEBRUARY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE: 22-APRIL-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/219,012
; FILING DATE: 28-MARCH-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,333
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; FILING DATE: 11-NOVEMBER-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/714,131
; FILING DATE: 10-JUNE-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/536,428
; FILING DATE: 11-JUNE-1990
; ATTORNEY/AGENT INFORMATION:
; NAME: Barry J. Swanson
; REGISTRATION NUMBER: 33,215
; REFERENCE/DOCKET NUMBER: NEX07/PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (303) 793-3333
; TELEFAX: (303) 793-3433
; INFORMATION FOR SEQ ID NO: 268:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 60 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
;
US-08-687-421-268

alignment_scores:
  Quality: 40.00      Length: 19
  Ratio: 2.667       Gaps: 0
  Percent Similarity: 78.947   Percent Identity: 42.105

alignment_block:
  US-09-439-311-2 x US-08-687-421-268 ..

Align seg 1/1 to: US-08-687-421-268 from: 1 to: 60

204 ThseryValgYthrglyleuglyAlaleuAlaclutlleasnaAgas 220
||||: |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
4 ACCCGGAGGCGGCGTAGCGTGTGAGCGCTGGCCGATGTGGAGCGCAGCA 53

220 nAlaasp 222
|||||
54 CTCGGAT 60

seq_name: /cgn2_6/ptodata/2/lna/5A_COMB.seq:US-08-482-882-97

seq_documentation_block:
; Sequence 97, Application US/08482882
; Patent No. 3773218
; GENERAL INFORMATION:
; APPLICANT: Gallatin, W. Michael
; TITLE OF INVENTION: ICAV-Related Materials and Methods
; NUMBER OF SEQUENCES: 116
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 S. Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/482,882
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/286,754
; FILING DATE:
; APPLICATION NUMBER: US 08/102,852
; FILING DATE: 05-AUG-1993
; PRIOR APPLICATION DATA:
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; APPLICATION NUMBER: US 08/009,266
; FILING DATE: 22-JAN-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/894,061
; FILING DATE: 05-JUN-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/889,724
; FILING DATE: 26-MAY-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/827,689
; FILING DATE: 27-JAN-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 5773218and, Greta E.
; REGISTRATION NUMBER: 35,302
; REFERENCE/DOCKET NUMBER: 32178
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 97:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 47 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
;
US-08-482-882-97

alignment_scores:
  Quality: 39.00      Length: 12
  Ratio: 3.900       Gaps: 0
  Percent Similarity: 83.333   Percent Identity: 66.667

alignment_block:
  US-09-439-311-2 x US-08-482-882-97/rev ..

Align seg 1/1 to reverse of: US-08-482-882-97 from: 1 to: 47

169 ArgpHecluthrglyserGlnSerPheSerSercly 180
||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
4 AGGATGGAGACTGGGTCAGCACGATTGGAGATGGA 9

seq_name: /cgn2_6/ptodata/2/lna/5A_COMB.seq:US-08-483-389-97

seq_documentation_block:
; Sequence 97, Application US/08483389
; Patent No. 5811517
; GENERAL INFORMATION:
; APPLICANT: Gallatin, W. Michael
; TITLE OF INVENTION: ICAV-RELATED PROTEIN
; NUMBER OF SEQUENCES: 118
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 233 South Wacker Drive/6300 Sears Tower
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/483,389
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/102,852
; FILING DATE: 05-AUG-1993
; PRIOR APPLICATION DATA:
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APPLICATION NUMBER: US 08/009,266  
FILING DATE: 22-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/894,061  
FILING DATE: 05-JUN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/889,724  
FILING DATE: 26-MAY-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,689  
FILING DATE: 27-JAN-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Suh, Young J.  
REGISTRATION NUMBER: P-41,337  
REFERENCE/DOCKET NUMBER: 27866/32760  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 474-6300  
TELEFAX: (312) 474-0448  
TELEX: (312) 474-6600  
INFORMATION FOR SEQ ID NO: 97:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-483-389-97

alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
US-09-439-311-2 x US-08-483-389-97/rev ..

Align seg 1/1 to reverse of: US-08-483-389-97 from: 1 to: 47

169 Arpnegiuthrglyserclnserphesergergly 180  
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44 AGATGAGACTGGCTCAGCAGATTGGAGTGCA 9

seq\_name: /cgn2\_6/ptodata/2/lna/5B\_COMB.seq:US-08-487-113D-97

seq\_documentation\_block:

; Sequence 97, Application US/08487113D  
; Patent No. 5817822  
; GENERAL INFORMATION:  
; APPLICANT: Gallatin, W. Michael  
; TITLE OF INVENTION: ICAM-Related Materials and Methods  
; NUMBER OF SEQUENCES: 120  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun  
; STREET: 6300 Sears Tower, 233 South Wacker Drive  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: United States of America  
; ZIP: 60606-6402  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/487,113D  
; FILING DATE:  
; CLASSIFICATION: 424  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/286,754  
; FILING DATE: 05-AUG-1994  
; PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/102,852  
FILING DATE: 05-AUG-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,266  
FILING DATE: 22-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/894,061  
FILING DATE: 05-JUN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/889,724  
FILING DATE: 26-MAY-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,689  
FILING DATE: 27-JAN-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: No. 5837822and, Greta E.  
REGISTRATION NUMBER: 35,302  
REFERENCE/DOCKET NUMBER: 32744  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 474-6300  
TELEFAX: (312) 474-0448  
TELEX: 25-3856  
INFORMATION FOR SEQ ID NO: 97:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-487-113D-97

alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
US-09-439-311-2 x US-08-487-113D-97/rev ..

Align seg 1/1 to reverse of: US-08-487-113D-97 from: 1 to: 47

169 Arpnegiuthrglyserclnserphesergergly 180  
||||:||||||||| |||:|||||  
44 AGATGAGACTGGCTCAGCAGATTGGAGTGCA 9

seq\_name: /cgn2\_6/ptodata/2/lna/5B\_COMB.seq:US-08-473-503-97

seq\_documentation\_block:

; Sequence 97, Application US/08473503  
; Patent No. 5869262  
; GENERAL INFORMATION:  
; APPLICANT: Gallatin, W. Michael  
; TITLE OF INVENTION: ICAM-Related Materials and Methods  
; NUMBER OF SEQUENCES: 116  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun  
; STREET: 6300 Sears Tower, 233 S. Wacker Drive  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60606  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/473,503  
; FILING DATE: 07-JUN-1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/286,754  
FILING DATE: 05-AUG-1994  
APPLICATION NUMBER: US 08/102,852  
FILING DATE: 05-AUG-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,266  
FILING DATE: 22-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/894,061  
FILING DATE: 05-JUN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/889,724  
FILING DATE: 26-MAY-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,689  
FILING DATE: 27-JAN-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: NO. 5869262and, Greta E.  
REGISTRATION NUMBER: 35,302  
REFERENCE/DOCKET NUMBER: 32178  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 474-6300  
TELEFAX: (312) 474-0448  
TELEX: 25-3856  
INFORMATION FOR SEQ ID NO: 97:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-473-503-97

alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667

## alignment\_block:

US-09-439-311-2 x US-08-473-503-97/rev ..

Align seg 1/1 to reverse of: US-08-473-503-97 from: 1 to: 47

169 ArgphegiurhglySerClnSerPheSerSergly 180

||||:||||||||||||| |||:|||||  
44 AGATGAGAGACTGGCTCAGACAGATTGGAGTGA 9

seq\_name: /cgn2\_6/plodata/2/lna/5B\_COMB.seq:US-08-483-932-97

## seq\_documentation\_block:

; Sequence 97, Application US/08483932

; Patent No. 5880268

; GENERAL INFORMATION:

; APPLICANT: Gallatin, W. Michael

; APPLICANT: Vazeux, Rosemay

; TITLE OF INVENTION: ICAM-Related Materials and Methods

; NUMBER OF SEQUENCES: 116

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun

; STREET: 6300 Sears Tower, 233 S. Wacker Drive

; CITY: Chicago

; STATE: Illinois

; COUNTRY: USA

; ZIP: 60606

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/483,932

; FILING DATE: 07-JUN-1995

CLASSIFICATION: 530  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/286,754  
FILING DATE: 05-AUG-1994  
APPLICATION NUMBER: US 08/102,852  
FILING DATE: 05-AUG-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,266  
FILING DATE: 22-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/894,061  
FILING DATE: 05-JUN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/889,724  
FILING DATE: 26-MAY-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,689  
FILING DATE: 27-JAN-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: NO. 5880268and, Greta E.  
REGISTRATION NUMBER: 35,302  
REFERENCE/DOCKET NUMBER: 32178  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 474-6300  
TELEFAX: (312) 474-0448  
TELEX: 25-3856  
INFORMATION FOR SEQ ID NO: 97:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-483-932-97

alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667

## alignment\_block:

US-09-439-311-2 x US-08-483-932-97/rev ..

Align seg 1/1 to reverse of: US-08-483-932-97 from: 1 to: 47

169 ArgphegiurhglySerClnSerPheSerSergly 180

||||:||||||||||||| |||:|||||  
44 AGATGAGAGACTGGCTCAGACAGATTGGAGTGA 9







US-09-439-311-2 x AR153101 ..

Align seg 1/1 to: AR153101 from: 1 to: 30

97 GlnAspGlyGlnSerLeuYsrHrArGThr 106  
|||||  
1 CAAAGTGGCAGACAGATTAAAAACAGAACT 30

seq\_name: gb\_pat:A93365

seq\_documentation\_block:

LOCUS A93365 50 bp DNA PAT 22-JAN-2000  
DEFINITION Sequence 4 from Patent WO9744451.  
ACCESSION A93365  
VERSION A93365.1 GI:6741628

KEYWORDS

SOURCE unidentified.

ORGANISM unidentified.

REFERENCE 1 (bases 1 to 50)

AUTHORS Paesen,G.C. and Nuttall,P.A.  
TITLE VASOACTIVE AMINE BINDING MOLECULES  
JOURNAL Patent: WO 9744451-A 4 27-NOV-1997;  
OXFORD VACS LTD (GB); PASEN GUIDO CHRISTIAN (GB)

FEATURES

1. 50 Location/Qualifiers

/organism="unidentified"

/db\_xref="taxon:32644" 17 t

BASE COUNT 11 a 8 c 14 g 17 t

ORIGIN

alignment\_scores:

Quality: 41.00 Length: 10  
Ratio: 4.100 Gaps: 0  
Percent Similarity: 100.000 Percent Identity: 70.000

alignment\_block:

US-09-439-311-2 x A93365 ..

Align seg 1/1 to: A93365 from: 1 to: 50

262 SerAspGlyAspGlyAsnGlySerLeuIle 271  
|||||  
7 AGTGATGGTGATGATGATGATCCCTCTG 36

seq\_name: gb\_pat:A12323

seq\_documentation\_block:

LOCUS A12323 51 bp DNA PAT 06-DEC-1993  
DEFINITION Oligonucleotide.  
ACCESSION A12323  
VERSION A12323.1 GI:491330

KEYWORDS

SOURCE synthetic construct.

ORGANISM synthetic construct.

REFERENCE 1 (bases 1 to 51)

AUTHORS HYBRID PROTEINS OR POLYPEPTIDES  
TITLE Patent: WO 8802757-A 24 21-APR-1988;  
JOURNAL Location/Qualifiers

FEATURES

1. 51 /organism="synthetic construct"

/db\_xref="taxon:32630" 2 t

BASE COUNT 15 a 27 c 7 g 2 t

ORIGIN

alignment\_scores:

Quality: 41.00 Length: 17  
Ratio: 3.154 Gaps: 0  
Percent Similarity: 76.471 Percent Identity: 47.059

alignment\_block:

US-09-439-311-2 x A12323/rev ..

Align seg 1/1 to reverse of: A12323 from: 1 to: 51

300 GlyArgGlyIleLysIleThrGlySerIleGlyValGlyAlaGlyIleLe 316  
|||  
51 GGGATCGGGGTTGGCGTTGGCGTTGGCGTTGGCGTTGGCGATCCT 2

316 u 316

1 c 1

seq\_name: gb\_pat:A12324

seq\_documentation\_block:

LOCUS A12324 51 bp DNA PAT 06-DEC-1993  
DEFINITION Oligonucleotide.  
ACCESSION A12324  
VERSION A12324.1 GI:489519

KEYWORDS

SOURCE synthetic construct.

ORGANISM synthetic construct.

REFERENCE 1 (bases 1 to 51)

AUTHORS HYBRID PROTEINS OR POLYPEPTIDES  
TITLE Patent: WO 8802757-A 25 21-APR-1988;  
JOURNAL Location/Qualifiers

FEATURES

1. 51 /organism="synthetic construct"

/db\_xref="taxon:32630" 15 t

BASE COUNT 2 a 7 c 27 g 15 t

ORIGIN

alignment\_scores:

Quality: 41.00 Length: 17  
Ratio: 3.154 Gaps: 0  
Percent Similarity: 76.471 Percent Identity: 47.059

alignment\_block:

US-09-439-311-2 x A12324 ..

Align seg 1/1 to: A12324 from: 1 to: 51

300 GlyArgGlyIleLysIleThrGlySerIleGlyValGlyAlaGlyIleLe 316  
|||  
1 GGGATCGGGGTTGGCGTTGGCGTTGGCGTTGGCGTTGGCGATCCT 50

316 u 316

51 c 51

seq\_name: gb\_pat:A12596

seq\_documentation\_block:

LOCUS A12596 51 bp DNA PAT 05-JAN-1994  
DEFINITION Oligonucleotide.  
ACCESSION A12596  
VERSION A12596.1 GI:491421

KEYWORDS

SOURCE synthetic construct.

ORGANISM synthetic construct.

REFERENCE 1 (bases 1 to 51)

AUTHORS RECOMBINANT VIRUS  
TITLE Patent: WO 8701386-A 12 12-MAR-1987;  
JOURNAL Location/Qualifiers

FEATURES

1. 51 /organism="synthetic construct"

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BASE COUNT      15 a      27 c      7 g      2 t
ORIGIN
Alignment_scores:
    Quality:   41.00          Length:   17
    Ratio:     3.154         Gaps:       0
Percent Similarity: 76.471      Percent Identity: 47.059
Alignment_block:
US-09-439-311-2 x A12596/rev ..
Allign seg 1/1 to reverse of: A12596 from: 1 to: 51
300 glyargclyleylserlterhrglyserllleglyvalglalyalaglyilele 316
||| ||||: :: :|||:::|||||:|||||:|||||
51 GGGAATCGGGGTTGGCGTGTGGGGTGTGGCGTGTGGGGTGTGGCGTGTGGGATCT 2
316 u 316
1 C 1
seq_name: gb_pat:A12597
seq_documentation_block:
LOCUS A12597 51 bp DNA PAT 05-JAN-1994
DEFINITION oligonucleotide.
ACCESSION A12597
VERSION A12597.1 GI:489543
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial construct.
REFERENCE 1 (bases 1 to 51)
AUTHORS
TITLE RECOMBINANT VIRUS
JOURNAL Patent: WO 8701386-A 13 12-MAR-1987;
FEATURES
Location/Qualifiers
source 1..51
/organism="synthetic construct"
/db_xref="taxon:32630"
BASE COUNT 2 a 7 c 27 g 15 t
ORIGIN
Alignment_scores:
    Quality:   41.00          Length:   17
    Ratio:     3.154         Gaps:       0
Percent Similarity: 76.471      Percent Identity: 47.059
Alignment_block:
US-09-439-311-2 x A12597 ..
Allign seg 1/1 to: A12597 from: 1 to: 51
300 glyargclyleylserlterhrglyserllleglyvalglalyalaglyilele 316
||| ||||: :: :|||:::|||||:|||||:|||||
1 GGGAATCGGGGTTGGCGTGTGGGGTGTGGCGTGTGGGGTGTGGGATCT 50
316 u 316
1 C 1
seq_name: gb_pat:AX011428
seq_documentation_block:
LOCUS AX011428 59 bp DNA PAT 06-SEP-2000
DEFINITION Sequence 105 from Patent W0955907.
ACCESSION AX011428
VERSION AX011428.1 GI:9997978
KEYWORDS
synthetic construct.
```

```

ORGANISM      synthetic construct
              artificial sequence.
REFERENCE     1 (bases 1 to 59)
AUTHORS      Koetter,P., Entian,K.D. and Diu-Hercend,A.
TITLE        Method for screening antimycotic substances using essential genes
              from S. Cerevisiae
JOURNAL      Patent: WO 9955907-A 105 04-NOV-1999;
              KOETTER PETER (DE); EMTIAN KARL DIETER (DE); DIU HERCEND ANITA
              (FR); HOECHST MARION ROUSSEL INC (FR)
FEATURES
  source      1..59
              /organism="synthetic construct"
              /db_xref="taxon:32630"
              /note="primer YMR290c-S1"
BASE COUNT   8 a      12 c      15 g      24 t
ORIGIN
alignment_scores:
  Quality:    40.00      Length:      15
  Ratio:      3.077      Gaps:        0
Percent Similarity: 86.667      Percent Identity: 53.333

alignment_block:
US-09-439-311-2 x AX011428      ..

Align seg 1/1 to: AX011428 from: 1 to: 59

163 SerLysIleGlyValThrArgPheGluThrGlySerGlnSerPhe 177
      ::::::::::::::: ||::::::::::::: ||::::
9  ACGTCGTTGTGGTATTGGCGTTTTCACACTGGCAGCTGACGCTC 53

seq_name: gb_pat:ARI25926

seq_documentation_block:
LOCUS      ARI25926      60 bp      DNA      PAT      16-MAY-2001
DEFINITION Sequence 268 from patent US 6177557.
VERSION     ARI25926
KEYWORDS    ARI25926.1 GI:14111988

SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 60)
AUTHORS     Janjic,N., Gold,L. and Tasset,D.
TITLE       High affinity ligands of basic fibroblast growth factor and
              thrombin
JOURNAL     Patent: US 6177557-A 268 23-JAN-2001;
              Location/Qualifiers
FEATURES
  source      1..60
              /organism="unknown"
BASE COUNT   10 a      11 c      29 g      10 t
ORIGIN
alignment_scores:
  Quality:    40.00      Length:      19
  Ratio:      2.667      Gaps:        0
Percent Similarity: 78.947      Percent Identity: 42.105

alignment_block:
US-09-439-311-2 x ARI25926      ..

Align seg 1/1 to: ARI25926 from: 1 to: 60

204 ThrSerValGlyThrGlyLeuGlyAlaLeuAlaGluGluIleAsnArgAs 220
      ::::::::::: ||::::::::::::: ||::::::::::::: ::
4  ACCGGGAGGAGGCGGTGAGCGCTTGGAGCGCTTGGCCATGTGTAGGACACGA 53

220 nlaIaSp 222
      ::::::::::
54 CTCGGAT 60

```

Align seg 1/1 to reverse of: AR013897 from: 1 to: 47

Align seg 1/1 to reverse of: AR042511 from: 1 to: 47

169 ArpPhegluFrhGlySerGlnSerPheSerSergly 180  
 |||::|||  
 44 AGGATGGAGACTGGGTCTCAGCACGATTTGGGAGTGA 9

seq\_name: gb\_pat:AR058391

seq\_documentation\_block:

LOCUS AR058391 47 bp DNA PAT 29-SEP-1999  
 DEFINITION Sequence 97 from patent US 5837822.  
 ACCESSION AR058391  
 VERSION AR058391.1 GI:5983968  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Gallatin,W.Michael and Vazeux,R.  
 TITLE Humanized antibodies specific for ICAM related protein  
 JOURNAL Patent: US 5837822-A 97 17-NOV-1998;  
 FEATURES  
 source 1..47  
 /organism="unknown"  
 BASE COUNT 9 a 21 c 7 g 10 t  
 ORIGIN

alignment\_scores:

Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:

US-09-439-311-2 x AR058391/rev ..

Align seg 1/1 to reverse of: AR058391 from: 1 to: 47

169 ArpPhegluFrhGlySerGlnSerPheSerSergly 180  
 |||::|||  
 44 AGGATGGAGACTGGGTCTCAGCACGATTTGGGAGTGA 9

seq\_name: gb\_pat:AR08217

seq\_documentation\_block:

LOCUS AR08217 47 bp DNA PAT 07-SEP-2000  
 DEFINITION Sequence 97 from patent US 5989843.  
 ACCESSION AR08217  
 VERSION AR08217.1 GI:10014980  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 47)  
 AUTHORS Gallatin,W.Michael and Vazeux,R.  
 TITLE Methods for identifying modulators of protein kinase C  
 JOURNAL Patent: US 5989843-A 97 23-NOV-1999;  
 FEATURES  
 source 1..47  
 /organism="unknown"  
 BASE COUNT 9 a 21 c 7 g 10 t  
 ORIGIN

alignment\_scores:

Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:

US-09-439-311-2 x AR08217/rev ..

Align seg 1/1 to reverse of: AR08217 from: 1 to: 47

169 ArpPhegluFrhGlySerGlnSerPheSerSergly 180  
 |||::|||  
 44 AGGATGGAGACTGGGTCTCAGCACGATTTGGGAGTGA 9

Wed Apr 17 07:36:47 2002

us-09-439-311-2.rge

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GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

Run on: April 16, 2002, 23:26:24 ; Search time 1531.8 Seconds  
(without alignments)  
10759.030 Million cell updates/sec

Title: US-09-439-311-1

Perfect score: 999  
Sequence: 1 attacacaaatgttcacg.....ttaaaatggtgtagagat 999

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 1472140 seqs, 8248589755 residues

Total number of hits satisfying chosen parameters: 586436

Minimum DB seq length: 0  
Maximum DB seq length: 60

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenBml:\*

- 1: gb\_ba:\*
- 2: gb\_hlg:\*
- 3: gb\_in:\*
- 4: gb\_com:\*
- 5: gb\_ov:\*
- 6: gb\_pat:\*
- 7: gb\_ph:\*
- 8: gb\_pl:\*
- 9: gb\_pr:\*
- 10: gb\_ro:\*
- 11: gb\_sts:\*
- 12: gb\_sy:\*
- 13: gb\_un:\*
- 14: gb\_vl:\*
- 15: em\_ba:\*
- 16: em\_fun:\*
- 17: em\_hum:\*
- 18: em\_in:\*
- 19: em\_om:\*
- 20: em\_or:\*
- 21: em\_ov:\*
- 22: em\_pat:\*
- 23: em\_ph:\*
- 24: em\_pl:\*
- 25: em\_ro:\*
- 26: em\_sts:\*
- 27: em\_sy:\*
- 28: em\_un:\*
- 29: em\_vl:\*
- 30: em\_hlg\_hum:\*
- 31: em\_hlg\_inv:\*
- 32: em\_hlg\_rod:\*
- 33: em\_hlg\_hum:\*
- 34: em\_hlg\_inv:\*
- 35: em\_hlg\_rod:\*
- 36: em\_hlg\_other:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	26.8	2.7	30	6	ARI53100	ARI53100 Sequence
2	26.8	2.7	30	6	ARI53101	ARI53101 Sequence
3	21.4	2.1	60	3	AF320167	AF320167 Drosophila
C 4	20.8	2.1	24	6	AR001228	AR001228 Sequence
C 5	20.8	2.1	24	6	AR008251	AR008251 Sequence
C 6	20.8	2.1	24	6	AR010178	AR010178 Sequence
C 7	20.8	2.1	24	6	AR064102	AR064102 Sequence
C 8	20.8	2.1	24	6	I38291	I38291 Sequence 6
C 9	20.8	2.1	44	6	AX052949	AX052949 Sequence
C 10	20.8	2.1	58	6	AR061278	AR061278 Sequence
C 11	20.6	2.1	60	6	AB0405	AB0405 Sequence 17
12	20.4	2.0	51	6	AX160493	AX160493 Sequence
13	20.4	2.0	51	6	AX160494	AX160494 Sequence
14	20.4	2.0	57	3	AF320169	AF320169 Drosophila
15	20.4	2.0	57	3	AF320170	AF320170 Drosophila
16	20.4	2.0	57	3	AF320171	AF320171 Drosophila
17	20.4	2.0	57	3	AF320172	AF320172 Drosophila
18	20.4	2.0	57	3	AF320173	AF320173 Drosophila
19	20.4	2.0	57	3	AF320174	AF320174 Drosophila
20	20.4	2.0	57	3	AF320175	AF320175 Drosophila
21	20.4	2.0	57	3	AF320176	AF320176 Drosophila
22	20.4	2.0	57	3	AF320177	AF320177 Drosophila
23	20.4	2.0	57	3	AF320178	AF320178 Drosophila
24	20.4	2.0	57	3	AF320179	AF320179 Drosophila
25	20.2	2.0	52	6	A69988	A69988 Sequence 19
C 26	20	2.0	50	6	A17107	A17107 Oligonucleo
C 27	20	2.0	50	6	AR027492	AR027492 Sequence
C 28	20	2.0	57	6	AR077133	AR077133 Sequence
29	20	2.0	57	6	AR102800	AR102800 Sequence
30	20	2.0	57	6	I21136	I21136 Sequence 4
31	20	2.0	60	6	A38623	A38623 Sequence 16
32	20	2.0	60	6	AR040717	AR040717 Sequence
C 33	19.8	2.0	51	6	AX165291	AX165291 Sequence
C 34	19.8	2.0	59	6	AX011435	AX011435 Sequence
C 35	19.6	2.0	29	6	A20557	A20557 Oligonucleo
C 36	19.6	2.0	43	6	I11627	I11627 Sequence 12
C 37	19.6	2.0	51	6	AX160235	AX160235 Sequence
38	19.6	2.0	55	14	HIV1045041	U45041 Human Immun
39	19.6	2.0	57	6	A32988	A32988 Synthetic P
C 40	19.6	2.0	58	6	AR011305	AR011305 Sequence
C 41	19.6	2.0	58	6	I17943	I17943 Sequence 17
C 42	19.4	1.9	37	6	AR063204	AR063204 Sequence
43	19.4	1.9	49	14	ALAM03	V00054 Alfalfa mos
44	19.4	1.9	51	6	AX117493	AX117493 Sequence
C 45	19.4	1.9	51	6	AX162140	AX162140 Sequence

## ALIGNMENTS

RESULT 1  
ARI53100/c  
LOCUS ARI53100 30 bp DNA  
DEFINITION Sequence 102 from patent US 6235480.  
ACCESSION ARI53100  
VERSION ARI53100.1 GI:15120632  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)  
Shultz, J. William, Lewis, M. K., Laipe, D., Mandrekar, M., Kephart, D.,  
Rhodes, R. Byron, Andrews, C. Ann, Hartnett, J. Robert, Gu, T.,  
Olson, R. J., Wood, K. V. and Welch, R.  
Detection of nucleic acid hybrids  
Patent: US 6235480-A 102 22-MAY-2001;

TITLE JOURNAL  
FEATURES Location/Qualifiers  
1..30  
/organism="unknown"

BASE COUNT 5 a 5 c 4 g 16 t





JOURNAL Patent: US 5753444-A 6 19-MAY-1998;

FEATURES Location/Qualifiers  
Source 1..24  
/organism="unknown"

BASE COUNT 6 a 5 c 5 g 8 t

ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 24;

Best Local Similarity 91.7%; Pred. No. 1.9e+06; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 2;

QY 91 ggcctagaatcaactccgcagca 114  
|||||

Db 24 GGTCTTAGAATTAACCTCAGCAGCA 1

RESULT 6

LOCUS AR010178/c

DEFINITION Sequence 6 from patent US 5756701.

ACCESSION AR010178

VERSION AR010178.1 GI:3968983

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 24)

AUTHORS Wu, L., Coombs, J., Malmstrom, S.L. and Glass, M.J.

TITLE Specific oligonucleotide primer pairs and probes for discriminating

JOURNAL Patent: US 5756701-A 6 26-MAY-1998;

FEATURES Location/Qualifiers

source 1..24

BASE COUNT 6 a 5 c 5 g 8 t

ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 24;

Best Local Similarity 91.7%; Pred. No. 1.9e+06; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 2;

QY 91 ggcctagaatcaactccgcagca 114  
|||||

Db 24 GGTCTTAGAATTAACCTCAGCAGCA 1

RESULT 7

LOCUS AR064102/c

DEFINITION Sequence 6 from patent US 5846783.

ACCESSION AR064102

VERSION AR064102.1 GI:5993410

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 24)

AUTHORS Wu, L., Coombs, J., Malmstrom, S.L. and Glass, M.J.

TITLE Methods and apparatus for preparing, amplifying, and discriminating

JOURNAL Patent: US 5846783-A 6 08-DEC-1998;

FEATURES Location/Qualifiers

source 1..24

BASE COUNT 6 a 5 c 5 g 8 t

ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 24;

Best Local Similarity 91.7%; Pred. No. 1.9e+06; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 2;

QY 91 ggcctagaatcaactccgcagca 114  
|||||

Db 24 GGTCTTAGAATTAACCTCAGCAGCA 1

RESULT 8

LOCUS I38291/c

DEFINITION Sequence 6 from patent US 5612473.

ACCESSION I38291

VERSION I38291.1 GI:2086281

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 24)

AUTHORS Wu, L., Coombs, J., Malmstrom, S.L. and Glass, M.J.

TITLE Methods, kits and solutions for preparing sample material for

JOURNAL Patent: US 5612473-A 6 18-MAR-1997;

FEATURES Location/Qualifiers

source 1..24

BASE COUNT 6 a 5 c 5 g 8 t

ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 24;

Best Local Similarity 91.7%; Pred. No. 1.9e+06; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 2;

QY 91 ggcctagaatcaactccgcagca 114  
|||||

Db 24 GGTCTTAGAATTAACCTCAGCAGCA 1

RESULT 9

LOCUS AX052949

DEFINITION Sequence 55 from Patent WO0071755.

ACCESSION AX052949

VERSION AX052949.1 GI:12227051

KEYWORDS

SOURCE Synthetic construct.

ORGANISM Synthetic construct.

REFERENCE 1 (bases 1 to 44)

AUTHORS Kwoh, J.G., Macklin, J.J., Mitsis, P.G. and Ulmer, K.M.

TITLE Method for sequencing and characterizing polymeric biomolecules using

JOURNAL aptamers and a method for producing aptamers

Patent: WO 0071755-A 55 30-NOV-2000;

FEATURES Incorporated (US)

source 1..44

BASE COUNT 9 a 6 c 19 g 10 t

ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 44;

Best Local Similarity 70.0%; Pred. No. 1.8e+06; Indels 0; Gaps 0;

Matches 28; Conservative 0; Mismatches 12;

QY 747 tgcatacatggggttgtagtgaagtgcattatca 786  
|||||

Db 4 TGACACCACTGGGGTGGGTATGGGTAGCGTTTGGAATCA 43

RESULT 10

LOCUS AR061278/c

LOCUS AR061278 58 bp DNA PAT 29-SEP-1999  
DEFINITION Sequence 7 from patent US 5843650.  
ACCESSION AR061278  
VERSION AR061278.1 GI:5988969  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 58)  
AUTHORS Segev,D.  
TITLE Nucleic acid detection and amplification by chemical linkage of  
JOURNAL oligonucleotides  
FEATURES Patent: US 5843650-A 7 01-DEC-1998;  
source Location/Qualifiers  
1..58  
BASE COUNT 15 a 18 c 13 g 12 t  
ORIGIN

Query Match 2.1%; Score 20.8; DB 6; Length 58;  
Best Local Similarity 60.7%; Pred. No. 1.8e+06;  
Matches 34; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

Oy 789 tggatgatgagaatggtcttcaattcagctatcaatgctgtaaaagatacaactg 844  
Db 58 TGATGCTGAGAGATGGCCTCGCTTCATGCTGCCATGCAGACATGTTACACATG 3

RESULT 11  
LOCUS A80405 60 bp DNA PAT 21-JAN-2000  
DEFINITION Sequence 17 from Patent WO951771.  
ACCESSION A80405  
VERSION A80405.1 GI:6731293  
KEYWORDS  
SOURCE Archaeoglobus fulgidus.  
ORGANISM Archaeoglobus fulgidus.  
REFERENCE 1 (bases 1 to 60)  
AUTHORS Jensen,R. and Schouls,L.M.  
TITLE A METHOD OF INTERSTRAIN DIFFERENTIATION OF BACTERIA  
JOURNAL Patent: WO 951771-A 17 14-OCT-1999; JANSSEN RUDOLPH (NL)  
EMBLEN JOHANNES DIRK ANTHONIE (NL); JANSSEN RUDOLPH (NL)  
FEATURES source Location/Qualifiers  
1..60  
misc\_feature /db\_xref="taxon:2234"  
BASE COUNT 20 a 11 c 9 g 20 t  
ORIGIN

Query Match 2.1%; Score 20.6; DB 6; Length 60;  
Best Local Similarity 62.7%; Pred. No. 2e+06;  
Matches 32; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

Oy 561 aaactacacggtatcgaagaatttaattgatagtagtgatcttctac 611  
Db 10 AGACCAAAAGGAGTGAATCTTCAATCCCATTTTGGCTGATTTCAAC 60

RESULT 12  
LOCUS AX160493 51 bp DNA PAT 22-JUN-2001  
DEFINITION Sequence 3821 from Patent WO0140521.  
ACCESSION AX160493  
VERSION AX160493.1 GI:14541824  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 51)  
AUTHORS Shimkets,R.A. and Leach,M.  
TITLE Nucleic acids containing single nucleotide polymorphisms and  
JOURNAL methods of use thereof  
FEATURES Patent: WO 0140521-A 3821 07-JUN-2001;  
source Curagen Corporation (US)  
1..51  
misc\_feature Location/Qualifiers  
26 /db\_xref="taxon:9606"  
/note="1 of 2 allelic variants (3822 is other entry)"  
BASE COUNT 19 a 5 c 8 g 19 t  
ORIGIN

Query Match 2.0%; Score 20.4; DB 6; Length 51;  
Best Local Similarity 65.2%; Pred. No. 2.2e+06;  
Matches 30; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

Oy 858 taaagatgaaaatgtaactgttcttacttcggccgatgtaga 903  
Db 4 TAAAGATGAACAGTTAAGCCATTTTGTGGAAGATGTAGA 49

RESULT 13  
LOCUS AX160494 51 bp DNA PAT 22-JUN-2001  
DEFINITION Sequence 3822 from Patent WO0140521.  
ACCESSION AX160494  
VERSION AX160494.1 GI:14541825  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 51)  
AUTHORS Shimkets,R.A. and Leach,M.  
TITLE Nucleic acids containing single nucleotide polymorphisms and  
JOURNAL methods of use thereof  
FEATURES Patent: WO 0140521-A 3822 07-JUN-2001;  
source Curagen Corporation (US)  
1..51  
misc\_feature Location/Qualifiers  
26 /db\_xref="taxon:9606"  
/note="2 of 2 allelic variants (3821 is other entry)"  
BASE COUNT 18 a 5 c 8 g 20 t  
ORIGIN

Query Match 2.0%; Score 20.4; DB 6; Length 51;  
Best Local Similarity 65.2%; Pred. No. 2.2e+06;  
Matches 30; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

Oy 858 taaagatgaaaatgtaactgttcttacttcggccgatgtaga 903  
Db 4 TAAAGATGAACAGTTAAGCCATTTTGTGGAAGATGTAGA 49

RESULT 14  
LOCUS AF320169 57 bp DNA INV 23-APR-2001  
DEFINITION Sequence 3822 from Patent WO0140521.  
ACCESSION AF320169  
VERSION AF320169.1 GI:13752326  
KEYWORDS  
SOURCE Drosophila pseudoobscura.  
ORGANISM Drosophila pseudoobscura

REFERENCE  
AUTHORS  
TITLE

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.  
1 (bases 1 to 57)  
Noor, M.A., Kilman, R.M. and Machado, C.A.  
Evolutionary history of microsatellites in the obscure group of

JOURNAL  
MOL. BIOL. EVOL. 18 (4), 551-556 (2001)  
MEDLINE  
21165327

PUBMED	11264406
REFERENCE	2 (bases 1 to 57)

TITLE	Author(s)
Direct Submission	MOORE/RENNETT / ALTMAN, R. and MACHUGO, C. R.

Rd., Piscataway, NJ 08854, USA

source  
1. :57  
/organism="Prosophi

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/db_xref="taxon:7237"
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/ gene="bcd"

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/gene="bcd"
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Best Local Similarity      61.1%: Pred. No. 2.2e+06;
Matches 33; Conservative 0; Mismatches 21; Indels 0; Gaps 0

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Oy      89 caggtccttagaatcaactccgcagcagatgatgcttcagygatgycgataagcag 142
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Db      4  CAGTTCCTTTCAGACACACAGCAGCAGCAGCTTCATCAGCAGCAGCAGCAACAGCAG 57

```

RESULT 15  
AE220170

0170763W

LOCUS	AF320170	57 bp	DNA	INV	23-APR-2001
DEFINITION	Drosophila pseudoobscura strain Mather17 b1coid (bcd) gene, partial				

ACCESSION	AF320170
VERSION	AF320170.1
KEYWORDS	GI:13752328

NETWORKS  
 SOURCE  
 ORGANISM

REFERENCE AUTHORS TITLE
1 (bases 1 to 57) Noor, M.A., Kliman, R.M. and Machado, C.A. Evolutionary history of microsatellites in the obscure group of

*Drosophila*  
MOL. BIOL. EVOL. 18 (4), 551-556 (2001)  
JOURNAL  
MEDIANE 21165327

MEDLINE	21105327
PUBMED	11264406
REFERENCE	2 (bases 1 to 571)

**AUTHORS** NOOR, M.A.F., Klíman, R.M. and Machado, C.A.  
**TITLE** Direct Submission

**JOURNAL** Submitted (08-NOV-2000) Genetics, Rutgers University, 604 Allison Rd., Piscataway, NJ 08854, USA

FEATURES	Location/Qualifiers
source	1. .57

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/db xref="taxon:7237"

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/gene acc
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/protein_id="AAK38622.1"
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COUNT      19 a      19 c      12 a      7 t      :
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Query Match	2.0%;	Score 20.4;	DB 3;	Length 57;
Best Local Similarity	61.1%;	Pred. No. 2.2e+06;		
Matches 33; Conservative	0;	Mismatches 21;	Indels 0;	Gaps 0;

**Qy** 89 cagtccttgaatcaactccgcagcagatgatgtcttcaggatgycgategcag 142  
||| ||| | ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||  
**Db** 4 CAGTCTCTTCAGACACAAGCAGCAGCAGCTCCATCAGCAGCAACAAGCAG 57

Search completed: April 17, 2002, 01:49:31  
Job time: 8587 sec



GenCore version 4.5  
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 17, 2002, 01:18:44 ; Search time 172.86 Seconds  
(without alignments)  
4954.691 Million cell updates/sec

Title: US-09-439-311-1

Perfect score: 999  
Sequence: 1 attacacaatgttcgacg.....ttaaaatgtagttagatg 999

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapept 1.0

Searched: 930621 seqs, 428662619 residues

Total number of hits satisfying chosen parameters: 1026190

Minimum DB seq length: 0  
Maximum DB seq length: 60

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :  
N.Geneseq\_1101:\*

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2:	/SID2/gcgdata/geneseq/geneseq/NA1981.DAT:*
3:	/SID2/gcgdata/geneseq/geneseq/NA1982.DAT:*
4:	/SID2/gcgdata/geneseq/geneseq/NA1983.DAT:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	26.8	2.7	30	21	AAA86891
2	26.8	2.7	30	21	AAA86892
3	26.8	2.7	30	21	AAA93188
4	26.8	2.7	30	21	AAA93190
5	21.4	2.2	50	20	AA552169
6	21.4	2.1	51	19	AAV04223
7	21.4	2.1	58	19	AAV04224
8	21.2	2.1	54	21	AAA97417
9	21	2.1	27	21	AAA27148
10	21	2.1	33	21	AAA27149
11	20.8	2.1	24	18	AA760508

C 12	20.8	2.1	24	19	AAV31446	Campylobacter nucl
C 13	20.8	2.1	24	19	AAV25942	Oligonucleotide PC
C 14	20.8	2.1	24	19	AAV20847	Campylobacter CF04
C 15	20.8	2.1	44	22	AA716711	dGMP-specific apta
C 16	20.8	2.1	50	18	AAV76100	Staphylococcus aur
C 17	20.8	2.1	51	22	AAH89291	Human coding sequ
C 18	20.8	2.1	58	19	AAV15200	Human serrate 2 PC
C 19	20.6	2.1	47	21	AAV67004	Human map-related
C 20	20.6	2.1	51	18	AA770191	Human map-related
C 21	20.6	2.1	51	20	AAV16831	Primer SEQ ID NO:2
C 22	20.4	2.0	47	21	AAV66354	Human delta-1 gene
C 23	20.4	2.0	47	21	AAZ68687	Human map-related
C 24	20.4	2.0	50	16	AAV25074	Human gene signatu
C 25	20.2	2.0	50	22	AAH25534	PCR primer used to
C 26	20.2	2.0	50	22	AAH25535	PCR primer used to
C 27	20.2	2.0	59	21	AAV24094	Human secreted pro
C 28	20.2	2.0	60	16	AAV21652	Human gene signatu
C 29	20	2.0	39	19	AAV05387	Primer PAS3BACREV
C 30	20	2.0	39	21	AAV55959	Sequencing and PCR
C 31	20	2.0	44	21	AAZ55398	Neisseria species
C 32	20	2.0	52	20	AAV16314	Human delta-2 prim
C 33	20	2.0	57	20	AAZ07541	ST receptor peptid
C 34	20	2.0	60	15	AAQ66366	McPC603 V-min gene
C 35	20	2.0	60	17	AAV38741	Moraxella outer me
C 36	19.8	2.0	45	22	AAV30242	Dystrophin exon 3
C 37	19.8	2.0	47	21	AAZ69525	Human map-related
C 38	19.8	2.0	59	21	AAZ96831	S. cerevisiae gene
C 39	19.6	2.0	36	14	AAQ40036	Sequence of revers
C 40	19.6	2.0	43	16	AAQ89640	Reverse primer 99R
C 41	19.6	2.0	47	21	AAZ66795	Human map-related
C 42	19.6	2.0	47	21	AAZ67933	Human map-related
C 43	19.6	2.0	57	13	AAQ23824	Primer Huc1ambd4FO
C 44	19.6	2.0	58	13	AAQ35636	SIV env primer SIV
C 45	19.6	2.0	58	14	AAQ35359	PCR primer STEVEN4

## ALIGNMENTS

RESULT 1	AAA86891/c	standard; DNA: 30 BP.
ID	AAA86891	
XX		
AC	AAA86891:	
XX		
DT	15-JAN-2001 (first entry)	
XX		
DE	Probe to Campylobacter jejuni.	
XX		
KW	Detection: nucleic acid hybrid; depolymerisation; analysis: SNP;	
KW	single nucleotide polymorphism; identification; viral load; probe;	
KW	genotyping; medical marker diagnostic; primer: target; mutation;	
KW	genetic disease: ss.	
XX		
OS	Campylobacter jejuni.	
XX		
PN	MO200049180-A1.	
PD	24-AUG-2000.	
XX		
PF	18-FEB-2000: 2000WO-US04242.	
XX		
PR	18-FEB-1999: 99US-0252436.	
PR	21-JUL-1999: 99US-0358972.	
PR	25-AUG-1999: 99US-0383316.	
XX		
PA	(PROM-) PROMEGA CORP.	
XX		
PI	Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;	
XX	Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;	
XX	WPI: 2000-565377/52.	

PT Determining presence or absence of a predetermined endogenous nucleic  
PT acid sequence by using an enzyme that depolymerizes the 3' end of an  
PT oligonucleotide probe hybridized to a target sequence to release  
PT identifier nucleotides -  
XX  
XX  
PS Example; Page 321; 3899p; English.  
XX  
CC The present invention describes a method (M1) for determining the  
CC presence or absence of a predetermined endogenous nucleic acid target  
CC sequence (ENAT). The method comprises hybridizing a probe having an  
CC identifier nucleotide (IN) with ENAT which is treated with an enzyme  
CC that depolymerizes the 3' end of hybridised NA to release the INs.  
CC M1 is used for determining the number of known sequence repeats present  
CC in a nucleic acid target sequence in a nucleic acid sample. The method  
CC is also useful for determining whether a nucleic acid target sequence in  
CC a sample is an allele from a homozygous or heterozygous locus. The  
CC method is also useful for detection of mutations, translocations and  
CC SNPs in nucleic acids (including those associated with genetic disease),  
CC determination of viral load, species identification, sample  
CC contamination, and analysis of forensic samples. AAA86791 to AAA87079  
CC and AAB12817 represent sequence which are used in the exemplification of  
CC the present invention.  
CC N.B. There is a discrepancy between the SEQ ID NO: and sequences given  
CC in the examples, and the SEQ ID NO: and sequences given in the sequence  
CC listing from the present invention.  
XX  
SQ Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;

Query Match 2.7%; Score 26.8; DB 21; Length 30;  
Best Local Similarity 93.3%; Pred. No. 1.3e+03;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatggtcaagcttaaaacaagaact 318  
||||||| ||||| ||||| ||||| |||||  
Db 30 CAAGATGACACAAAGTTAAAAACAAGAACT 1

RESULT 2  
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XX  
AC AAA86892:  
XX  
DT 15-JAN-2001 (first entry)  
XX  
DE Probe to Campylobacter jejuni.  
XX  
KM Detection; nucleic acid hybrid; depolymerisation; analysis; SNP;  
KM single nucleotide polymorphism; identification; viral load; probe;  
KM genotyping; medical marker diagnostic; primer; target; mutation;  
KM genetic disease; ss.  
XX  
OS Campylobacter jejuni.  
XX  
PN WO200049180-A1.  
XX  
PD 24-AUG-2000.  
XX  
PE 18-FEB-2000; 2000MO-US04242.  
XX  
PR 18-FEB-1999; 99US-0252436.  
PR 21-JUL-1999; 99US-0358972.  
PR 25-AUG-1999; 99US-0383316.  
XX  
PA (PROM-) PROMEGA CORP.  
XX  
PI Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
XX  
DR WPI; 2000-565377/52.  
XX  
PT Determining presence or absence of a predetermined endogenous nucleic

PT acid sequence by using an enzyme that depolymerizes the 3' end of an  
PT oligonucleotide probe hybridized to a target sequence to release  
PT identifier nucleotides -  
XX  
XX  
PS Example; Page 321; 3899p; English.  
XX  
CC The present invention describes a method (M1) for determining the  
CC presence or absence of a predetermined endogenous nucleic acid target  
CC sequence (ENAT). The method comprises hybridizing a probe having an  
CC identifier nucleotide (IN) with ENAT which is treated with an enzyme  
CC that depolymerizes the 3' end of hybridised NA to release the INs.  
CC M1 is used for determining the number of known sequence repeats present  
CC in a nucleic acid target sequence in a nucleic acid sample. The method  
CC is also useful for determining whether a nucleic acid target sequence in  
CC a sample is an allele from a homozygous or heterozygous locus. The  
CC method is also useful for detection of mutations, translocations and  
CC SNPs in nucleic acids (including those associated with genetic disease),  
CC determination of viral load, species identification, sample  
CC contamination, and analysis of forensic samples. AAA86791 to AAA87079  
CC and AAB12817 represent sequence which are used in the exemplification of  
CC the present invention.  
CC N.B. There is a discrepancy between the SEQ ID NO: and sequences given  
CC in the examples, and the SEQ ID NO: and sequences given in the sequence  
CC listing from the present invention.  
XX  
SQ Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;

Query Match 2.7%; Score 26.8; DB 21; Length 30;  
Best Local Similarity 93.3%; Pred. No. 1.3e+03;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatggtcaagcttaaaacaagaact 318  
||||||| ||||| ||||| ||||| |||||  
Db 1 caagatgacacaaagttaaaacaagaact 30

RESULT 3  
AAA93188/c  
ID AAA93188 standard; DNA; 30 BP.  
XX  
AC AAA93188:  
XX  
DT 11-JAN-2001 (first entry)  
XX  
DE Campylobacter jejuni interrogation probe 11451.  
XX  
KM Campylobacter jejuni; nucleic acid detection; genomic typing;  
KM mutation detection; viral load determination; species identification;  
KM forensic analysis; probe; ss.  
XX  
OS Campylobacter jejuni.  
XX  
PN WO200049179-A1.  
XX  
PD 24-AUG-2000.  
XX  
PE 18-FEB-2000; 2000MO-US04176.  
XX  
PR 18-FEB-1999; 99US-0252436.  
PR 21-JUL-1999; 99US-0358972.  
PR 27-SEP-1999; 99US-0406147.  
XX  
PA (PROM-) PROMEGA CORP.  
XX  
PI Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
XX  
DR WPI; 2000-549282/50.  
XX  
PT Detecting the presence of predetermined exogenous nucleic acid target  
PT sequence useful for e.g. genotyping, comprises depolymerizing the 3'  
PT end of an oligonucleotide probe hybridized to a nucleic acid target

PT sequence -  
XX  
PS Claim 47; Page 187; 230pp; English.  
XX  
CC The present sequence is an interrogation probe which was used to detect a  
CC segment of the genome of *Campylobacter jejuni*. This was performed as part  
CC of a method for determining the presence of a known exogenous nucleic  
CC acid target sequence in a nucleic acid sample. The method comprises  
CC admixing a treated sample with a depolymerizing enzyme which releases one  
CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
CC The method is used for assaying nucleic acids for a particular native or  
CC mutant sequence, and for genomic typing. It is useful for detecting  
CC mutations, translocations, and single nucleotide polymorphisms, .  
CC determination of viral load, species identification, detection of sample  
CC contamination, and analysis of forensic samples. Compared with previous  
CC methods of detecting nucleic acid hybrids, the new method has higher  
CC sensitivity without the need for radiochemicals or electrophoresis. It is  
CC quantitative, highly reproducible and can be automated. The method can  
CC reliably detect as few as 10 copies of a virus in a sample, and is  
CC capable of providing multiple analyses in a single assay (multiplex  
CC assay).  
XX  
SQ Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;  
XX  
Query Match 2.7%; Score 26.8; DB 21; Length 30;  
Best Local Similarity 93.3%; Pred. No. 1.3e+03;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
OY 289 caagatggtcaaacgttaaaacaagaact 318  
Db 30 CAAGATGACCAAGTTTAAAAACAAGACT 1  
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX  
RESULT 4  
AA93190  
ID AAA93190 standard; DNA: 30 BP.  
XX  
AC AAA93190;  
XX  
XX 11-JAN-2001 (first entry)  
XX  
DE *Campylobacter jejuni* Interrogation probe 11450.  
XX  
KM *Campylobacter jejuni*; nucleic acid detection; genomic typing;  
KM mutation detection; viral load determination; species identification;  
KM forensic analysis; probe: ss.  
XX  
OS *Campylobacter jejuni*.  
XX  
PN WO200049179-A1.  
XX  
PD 24-AUG-2000.  
XX  
PF 18-FEB-2000; 2000WO-US04176.  
XX  
PR 18-FEB-1999; 99US-0252436.  
PR 21-JUL-1999; 99US-0358972.  
PR 27-SEP-1999; 99US-0406147.  
XX  
PA (PROM-) PROMEGA CORP.  
XX  
PI Shultz JM, Lewis MK, Lelape D, Mandrekar M, Kephart D, Rhodes RB;  
PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
XX  
DR WPI; 2000-549282/50.  
XX  
PT Detecting the presence of predetermined exogenous nucleic acid target  
PT sequence useful for e.g. genotyping, comprises depolymerizing the 3'  
PT end of an oligonucleotide probe hybridized to a nucleic acid target  
PT sequence -  
XX  
PS Claim 47; Page 187; 230pp; English.

XX  
CC The present sequence is an interrogation probe which was used to detect a  
CC segment of the genome of *Campylobacter jejuni*. This was performed as part  
CC of a method for determining the presence of a known exogenous nucleic  
CC acid target sequence in a nucleic acid sample. The method comprises  
CC admixing a treated sample with a depolymerizing enzyme which releases one  
CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
CC The method is used for assaying nucleic acids for a particular native or  
CC mutant sequence, and for genomic typing. It is useful for detecting  
CC mutations, translocations, and single nucleotide polymorphisms,  
CC determination of viral load, species identification, detection of sample  
CC contamination, and analysis of forensic samples. Compared with previous  
CC methods of detecting nucleic acid hybrids, the new method has higher  
CC sensitivity without the need for radiochemicals or electrophoresis. It is  
CC quantitative, highly reproducible and can be automated. The method can  
CC reliably detect as few as 10 copies of a virus in a sample, and is  
CC capable of providing multiple analyses in a single assay (multiplex  
CC assay).  
XX  
SQ Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;  
XX  
Query Match 2.7%; Score 26.8; DB 21; Length 30;  
Best Local Similarity 93.3%; Pred. No. 1.3e+03;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
OY 289 caagatggtcaaacgttaaaacaagaact 318  
Db 1 caagatgacacaagtttaaaacaagaact 30  
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX  
RESULT 5  
AA52169/C  
ID AA52169 standard; DNA: 50 BP.  
XX  
AC AA52169;  
XX  
XX 18-JUN-1999 (first entry)  
XX  
DE Synthetic plasmid synlux4 construction oligonucleotide R53.  
XX  
KM DNA plasmid; lux A; lux B; *Vibrio fischeri*; luciferase; promoter;  
KM tng kanamycin/neomycin phosphotransferase; DNA synthesis;  
KM replication competent double-stranded polynucleotide; ss.  
XX  
OS Synthetic.  
XX  
PN WO9914318-A1.  
XX  
PD 25-MAR-1999.  
XX  
PF 16-SEP-1998; 98WO-US19312.  
XX  
PR 16-SEP-1997; 97US-0059017.  
XX  
PA (TEXA ) UNIV TEXAS SYSTEM.  
XX  
PI Evans GA;  
XX  
XX WPI; 1999-244029/20.  
XX  
DR  
XX  
PT Synthesis of replication competent double-stranded polynucleotides  
PT Example 4; Fig 5E; 135pp; English.  
XX  
PS  
XX  
CC AAX52021-212 represent oligonucleotide primers that were used to  
CC construct a synthetic DNA plasmid sequence synlux4, to demonstrate the  
CC method of the invention. Within the synlux4 sequence are included the  
CC sequences of lux A, lux B, the A and B components of the *Vibrio fischeri*  
CC luciferase sequence, positions of pUC19 including the origin of  
CC replication and replication stability sequences, and the promoter and  
CC coding sequence for tng kanamycin/neomycin phosphotransferase. The  
CC specification describes a method for the synthesis of replication

CC competent double-stranded polynucleotides. The method comprises  
 CC generating a first set of oligonucleotides corresponding to the plus  
 CC strand and a second set corresponding to the minus strand and  
 CC annealing. The method can be used for preparing polynucleotides  
 CC encoding sequences involved in a biochemical pathway. In particular,  
 CC they can be used to produce polynucleotides encoding enzymes,  
 CC e.g. hexokinase, phosphohexose isomerase, phosphofructokinase-1,  
 CC aldolase, triose phosphate isomerase, glyceraldehyde-3-phosphate  
 CC dehydrogenase, phosphoglycerate kinase, phosphoglycerate mutase,  
 CC enolase or pyruvate kinase. They can also be used for the preparation  
 CC of viral particles, artificial genomes and artificial genetic systems.  
 XX

SQ Sequence 50 BP; 19 A; 8 C; 4 G; 19 T; 0 other;

Query Match 2.1%; Score 21.8; DB 20; Length 50;  
 Best Local Similarity 70.7%; Pred. No. 2.3e+04;  
 Matches 29; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

OY 808 ttaattcagctatcaatgctgtaagaatcaactggtgt 848  
 Db 43 TCATGTCGCGATTTAAAGATGTAAGATATTATGTAT 3

RESULT 6  
 ID AAV04223 standard; DNA: 51 BP.

XX AAV04223;

XX 22-JUN-1998 (first entry)

DE Human cardiac troponin I/troponin C 3' PCR primer.

XX Troponin I; troponin C; immunoassay; assay; analysis; human;  
 KM cardiac muscle; skeletal muscle; injury; myocardial infarction;  
 KW diagnosis; HctnI; HctnC; PCR; primer; ss.

OS Synthetic.

OS Homo sapiens.

PN W09739132-A1.

XX 23-OCT-1997.

XX 14-APR-1997; 97WO-US06147.

XX 11-APR-1997; 97US-0833743.

XX 16-APR-1996; 96US-0015772.

XX (UYMI-) UNIV MIAMI.

PI Potter JD;

XX WPI; 1998-062676/06.

XX Immunassay of mammalian troponin using stable standard for

PT comparison - specifically acid-dialysed solution or its lyophilisate

XX used for diagnosis of cardiac or skeletal muscle damage

XX Example 3; Page 34; 94pp; English.

XX This 3' PCR primer was used in the amplification of human cardiac  
 CC troponin I (HctnI) cDNA. It is a complementary sequence encoding  
 CC the C-terminal 8 amino acids of HctnI followed by the N-terminal 8  
 CC amino acids of human cardiac troponin C (HctnC). It was used with  
 CC a vector-based 5' primer in the PCR amplification of HctnI plasmid  
 CC DNA. HctnC DNA was also amplified (see AAV04223), and the PCR  
 CC products were used to construct a polynucleotide (see AAV04225)  
 CC encoding a HctnI/HctnC fusion protein (see AAW41571). The addition  
 CC of the calcium binding protein HctnC to HctnI was made to provide  
 CC more favourable solubility properties. The fusion protein can be  
 CC used as a standard in novel assays of mammalian troponin.

XX SQ Sequence 51 BP; 8 A; 17 C; 9 G; 17 T; 0 other;

Query Match 2.1%; Score 21.4; DB 19; Length 51;  
 Best Local Similarity 71.8%; Pred. No. 2.9e+04;  
 Matches 28; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

OY 505 agattgaacaggttcacaagtttcttcaggcact 543  
 Db 12 agatgcatcattcctcctcaactttcttcggccct 50

RESULT 7  
 ID AAV04224/c  
 XX AAV04224 standard; DNA: 58 BP.

XX AAV04224;

XX 22-JUN-1998 (first entry)

DE Human cardiac troponin C 5' PCR primer.

XX Troponin C; troponin I; immunoassay; assay; analysis; human;  
 KW cardiac muscle; skeletal muscle; injury; myocardial infarction;  
 KW diagnosis; HctnI; HctnC; PCR; primer; ss.

OS Synthetic.

OS Homo sapiens.

PN W09739132-A1.

XX 23-OCT-1997.

XX 14-APR-1997; 97WO-US06147.

XX 11-APR-1997; 97US-0833743.

XX 16-APR-1996; 96US-0015772.

XX (UYMI-) UNIV MIAMI.

PI Potter JD;

XX WPI; 1998-062676/06.

XX Immunassay of mammalian troponin using stable standard for

PT comparison - specifically acid-dialysed solution or its lyophilisate

XX used for diagnosis of cardiac or skeletal muscle damage

XX Example 3; Page 34; 94pp; English.

XX This 5' PCR primer was used with a vector-based 3' primer in the  
 CC amplification of human cardiac troponin C (HctnC) DNA. Human  
 CC cardiac troponin I (HctnI) DNA was also amplified (see AAV04223) and  
 CC the PCR products were used to construct a polynucleotide (see  
 CC AAV04225) encoding a HctnI/HctnC fusion protein (see AAW41571). The  
 CC addition of the calcium binding protein HctnC to HctnI was made to  
 CC provide more favourable solubility properties. The fusion protein  
 CC can be used as a standard in novel assays of mammalian troponin.

SQ Sequence 58 BP; 18 A; 10 C; 20 G; 10 T; 0 other;

Query Match 2.1%; Score 21.4; DB 19; Length 58;  
 Best Local Similarity 71.8%; Pred. No. 3e+04;  
 Matches 28; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

OY 505 agattgaacaggttcacaagtttcttcaggcact 543  
 Db 39 AGATGTCATCAGTCCTCCTCAAACTTTCTTGGCGCCCT 1

RESULT 8



OS	Campylobacter coli.
XX	
FN	WO200027205-A1.
XX	
PD	18-MAY-2000.
XX	
PE	12-NOV-1999; 99WO-US27195.
XX	
PR	12-NOV-1998; 98US-0108114.
XX	
PA	(USAT ) US SEC.
XX	
EI	Guerry P, Trust T, Burg E, Lee L;
XX	
DR	WPI; 2000-376214/32.
XX	
PT	Campylobacter FlaA protein and coding sequence, useful in reducing
PP	Campylobacter intestinal colonization -
XX	
PS	Disclosure; Page 7; 43pp; English.
XX	
CC	The flaA gene encodes the major flagellin subunit of the Campylobacter
CC	coli flagellar filament. Part of the FlaA polypeptide may be fused with
CC	the maltose binding protein of Escherichia coli to make a recombinant
CC	protein. When this protein is introduced into a host an immunological
CC	response is triggered. Therefore the recombinant protein may be used as
CC	a vaccine to protect against C. coli intestinal colonisation and the
CC	d diarrhoea it causes. This vaccine system is useful as it can
CC	prevent the development of Guillain-Barre syndrome (GBS) which is seen
CC	with whole cell Campylobacter vaccines. The present sequence is the
CC	flaA-11 PCR primer that was used to amplify part of the flaA gene.
XX	
SQ	Sequence 27 BP; 12 A; 6 C; 3 G; 6 T; 0 other;
XX	
OY	Query Match 2.1%; Score 21; DB 21; Length 27;
	Best Local Similarity 100.0%; Pred. No. 3.1e+04;
	Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB	1 attaacacaatgttcagca 21       7 attaacacaatgttcagca 27
XX	
RESULT 10	
ID	AAA27149/G
XX	AAA27149 standard; DNA; 33 BP.
AC	
XX	AAA27149;
XX	
DT	11-SEP-2000 (first entry)
XX	
DE	Campylobacter coli flaA gene primer flaA-2.
XX	
KW	Flagellin; flaA; diarrhoea; Guillain-Barre syndrome;
RW	vaccine; GBS; PCR primer; ss.
XX	
OS	Campylobacter coli.
XX	
PN	WO200027205-A1.
XX	
PD	18-MAY-2000.
XX	
PE	12-NOV-1999; 99WO-US27195.
XX	
PR	12-NOV-1998; 98US-0108114.
XX	
PA	(USAT ) US SEC.
XX	
EI	Guerry P, Trust T, Burg E, Lee L;
XX	
DR	WPI; 2000-376214/32.

PT Campylobacter FlaA protein and coding sequence, useful in reducing  
 PT Campylobacter intestinal colonization  
 XX  
 PS Disclosure; Page 7; 43pp; English.  
 CC The flaA gene encodes the major flagellin subunit of the Campylobacter  
 CC coli flagellar filament. Part of the FlaA polypeptide may be fused with  
 CC the maltose binding protein of Escherichia coli to make a recombinant  
 CC protein. When this protein is introduced into a host an immunological  
 CC response is triggered. Therefore the recombinant protein may be used as  
 CC a vaccine to protect against C. coli intestinal colonization and the  
 CC diarrhoea it causes. This vaccine system is useful as it can  
 CC prevent the development of Guillain-Barre syndrome (GBS) which is seen  
 CC with whole cell Campylobacter vaccines. The present sequence is the  
 CC flaA-2 PCR primer that was used to amplify part of the flaA gene.  
 XX  
 SQ Sequence 33 BP; 10 A; 8 C; 1 G; 14 T; 0 other:

Query Match 2.1%; Score 21; DB 21; Length 33;  
 Best Local Similarity 100.0%; Pred. No. 3.3e+04;  
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 979 gttaaaatgtagtagat 999  
 Db 33 GTTAAATAATGATGCTAGAGAT 13

RESULT 11  
 AAT60508/C  
 ID AAT60508 standard; DNA; 24 BP.  
 XX  
 AC AAT60508;  
 XX  
 DT 10-JUN-1997 (first entry)  
 XX  
 DE Primer CFOAR.4.  
 XX  
 KW PCR; polymerase chain reaction; amplify; infection; forensic science;  
 XX infectious pathogen; genetic disorder; genetic variance; primer; ss.  
 OS Synthetic.  
 XX  
 PN US5612473-A.  
 XX  
 PD 18-MAR-1997.  
 XX  
 PF 16-JAN-1996; 96US-0587209.  
 XX  
 PR 16-JAN-1996; 96US-0587209.  
 XX  
 PA (GULL-) GULL LAB.  
 PI Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 XX WPI; 1997-192163/17.  
 DR  
 XX  
 PT Processing samples for amplification of nucleic acid target  
 PT sequences - using extraction buffer containing at least one  
 PT detergent and a salt composition of greater than 1 molar  
 PT concentration  
 XX  
 PS Example 3; Column 17; 21pp; English.  
 XX  
 CC AAT60503-T60514 represent amplification primers for DNA sequences  
 CC present in a sample processed by the method of the invention. The  
 CC processing method of the invention comprises obtaining a sample of  
 CC material potentially containing the target nucleic acid sequences, and  
 CC mixing the sample with an external buffer solution. The buffer solution  
 CC comprises two detergents, and at least one salt composition present in a  
 CC greater than 1 M concentration. The mixture is then centrifuged to obtain  
 CC a supernatant portion, which is then heated before being recentrifuged  
 CC to precipitate the proteins, and obtaining a second supernatant portion,

CC from which nucleic acids are precipitated. The isolated nucleic acids  
 CC are then dissolved. The method provides a rapid means of preparing a  
 CC sample for amplification so that multiple analyses can be detected and  
 CC differentiated within a relatively short time period (typically less  
 CC than 5 hours with the novel pre-processing step taking less than 5  
 CC minutes). Typical applications of nucleic acid amplification include  
 CC detection of infections in patients, foodstuffs and for  
 CC diagnostic/forensic or quality control purposes, to discriminate between  
 CC multiple potential infectious pathogens, to diagnose genetic disorders or  
 CC to identify genetic variances.  
 XX  
 SQ Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other:

Query Match 2.1%; Score 20.8; DB 18; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04;  
 Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 91 gttcttagaatcaactcgcagca 114  
 Db 24 GGTCTTAGAATTAACCTCAGCAGCA 1

RESULT 12  
 AAV31446/C  
 ID AAV31446 standard; DNA; 24 BP.  
 XX  
 AC AAV31446;  
 XX  
 DT 11-AUG-1998 (first entry)  
 XX  
 DE Campylobacter nucleic acid sequence amplifying primer CFOAR.  
 XX  
 KM Salmonella; microorganism; detection; multiple analyte; PCR primer;  
 XX Yersinia; Escherichia coli; Campylobacter; ss.  
 OS Synthetic.  
 XX  
 PN US5756701-A.  
 XX  
 PD 26-MAY-1998.  
 XX  
 PF 06-AUG-1996; 96US-0692725.  
 XX  
 PR 16-JAN-1996; 96US-0587209.  
 XX  
 PR 06-AUG-1996; 96US-0692725.  
 XX  
 PA (GULL-) GULL LAB INC.  
 PI Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 XX WPI; 1998-321634/28.  
 DR  
 XX  
 PT Nucleic acid probes and primers - for detecting Salmonella, Yersinia  
 PT or E. coli  
 XX  
 PS Claim 5; Column 17; 21pp; English.  
 XX  
 CC This primer is used for the PCR amplification of Campylobacter nucleic  
 CC acid sequences. The invention provides nucleic acid probes and primers  
 CC for detecting Salmonella, Yersinia or E. coli. It provides methods and  
 CC apparatus for detecting and discriminating multiple analytes within a  
 CC test sample. The methods are simple, user-friendly, cost effective and  
 CC fast. The methods and the probes and primer sequences are used for  
 CC detecting the corresponding microorganisms in clinical samples.  
 XX  
 SQ Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other:

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04;  
 Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 91 ggtcttagaactcaactcgcagca 114  
 |||||  
 Db 24 GGTCTTAGAATTAACTCAGCAGCA 1

RESULT 13  
 AAV25942/c  
 ID AAV25942 standard; DNA: 24 BP.  
 XX  
 AC AAV25942;  
 XX  
 DT 15-JUL-1998 (first entry)  
 XX  
 DE Oligonucleotide PCR primer CFO4R gene.  
 XX  
 KM Sequence-specific probe: enterohaemorrhagic; *Escherichia coli*;  
 KM *Salmonella*; *Campylobacter*; *Shigella*; *Yersinia*; beta-globin;  
 KM *gastroenteritis*; PCR primer: ss.  
 XX  
 OS Synthetic.  
 OS *Campylobacter* sp.  
 XX  
 PN US5753444-A.  
 PD 19-MAY-1998.  
 PF 07-AUG-1996; 96US-0689235.  
 XX  
 PR 16-JAN-1996; 96US-0587209.  
 PR 07-AUG-1996; 96US-0689235.  
 XX  
 PA (GULL-) GULL LAB INC.  
 XX  
 PI Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 DR WPI; 1998-311393/27.  
 XX  
 PT Distinguishing between similar nucleic acid samples - using  
 PT sequence-specific probes e.g. between enterohaemorrhagic and normal  
 PT *Escherichia coli*  
 XX  
 PS Example 3; Column 17; 21pp; English.  
 CC The present sequence represents a PCR primer used in an example of the  
 CC present invention. The present invention describes a method for  
 CC detecting mismatches between first and second nucleic acid sequences  
 CC having at least one base difference. The method comprises: (a) obtaining  
 CC at least one labelled probe consisting of an oligonucleotide sequence  
 CC spanning the location of at least one base difference between the first  
 CC and second sequences, where the oligonucleotide sequence contains at  
 CC least one neutral base molecule in a position other than the position of  
 CC the base difference(s) but is otherwise exactly complementary to the  
 CC first sequence, so that the probe hybridises more weakly with the second  
 CC sequence than with the first sequence; (b) mixing the probe(s) with the  
 CC first and second sequences under hybridisation conditions; (c)  
 CC dissociating any probe/second sequence hybrids; and (d) detecting  
 CC probe/first sequence hybrids. The method can be used to distinguish  
 CC between similar DNA/RNA sequences in a sample, especially to distinguish  
 CC enterohaemorrhagic *E. coli* O157:H7 from other *E. coli* strains e.g. in  
 CC stool samples from people suffering from gastroenteritis, caused  
 CC specifically by enterohaemorrhagic *E. coli*. Use of the method shortens  
 CC the time between sample preparation to obtaining results, than has been  
 CC possible with previous similar procedures.  
 XX  
 SO Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04;  
 Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 OY 91 ggtcttagaactcaactcgcagca 114

Db 24 GGTCTTAGAATTAACTCAGCAGCA 1  
 |||||

RESULT 14  
 AAV20847/c  
 ID AAV20847 standard; DNA: 24 BP.  
 XX  
 AC AAV20847;  
 XX  
 DT 01-JUL-1998 (first entry)  
 XX  
 DE *Campylobacter* CFO4R gene PCR primer.  
 XX  
 KM *Escherichia coli* strain O157:H7; detection; microorganism; infection;  
 KM enterohaemorrhagic; PCR primer: ss.  
 XX  
 OS Synthetic.  
 OS *Campylobacter* sp.  
 XX  
 PN US5738995-A.  
 PD 14-APR-1998.  
 PF 07-AUG-1996; 96US-0689236.  
 XX  
 PR 16-JAN-1996; 96US-0587209.  
 PR 07-AUG-1996; 96US-0689236.  
 XX  
 PA (GULL-) GULL LAB INC.  
 XX  
 PI Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 DR WPI; 1998-26031/23.  
 XX  
 PT Probes for detecting *Escherichia coli* strain O157:H7 - useful for  
 PT diagnosis of enterohaemorrhagic *Escherichia coli* infection(s)  
 XX  
 PS Example 3; Column 17; 21pp; English.  
 CC The present sequence represents a PCR primer used in an example of the  
 CC present invention. The present invention describes probes used in the  
 CC detection of *Escherichia coli* strain O157:H7 in a sample. The method of  
 CC detection comprises: (a) obtaining at least 1 probe specifically given  
 CC in the specification, labelled with a label that permits probe detection  
 CC when hybridised to a complementary nucleic acid sequence which is  
 CC specific for a nucleic acid sequence of the microorganism; (b)  
 CC hybridising the probes and the sample, and (c) detecting hybrids  
 CC comprising the probes and the nucleic acid sequences. The method and  
 CC probes may be used for diagnosis of enterohaemorrhagic *E. coli*  
 CC infections. The methods and the materials permit the detection and  
 CC discrimination of multiple analyses.  
 XX  
 SO Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04;  
 Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 OY 91 ggtcttagaactcaactcgcagca 114  
 |||||  
 Db 24 GGTCTTAGAATTAACTCAGCAGCA 1

RESULT 15  
 AAF16711  
 ID AAF16711 standard; DNA: 44 BP.  
 XX  
 AC AAF16711;  
 DT 09-MAR-2001 (first entry)



GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

Run on: April 17, 2002, 01:23:59 ; Search time 86.97 Seconds  
(without alignments)  
2601.489 Million cell updates/sec

Title: US-09-439-311-1

Perfect score: 999

Sequence: 1 attacacaaatgtgtcagc.....ttaaaatgatgtagatagat 999

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 351203 seqs, 113238999 residues

Total number of hits satisfying chosen parameters: 515962

Minimum DB seq length: 0  
Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued\_Patents\_NA:\*  
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2: /cgn2\_6/prodata/2/lna/5B.COMB.seq:\*  
3: /cgn2\_6/prodata/2/lna/6A.COMB.seq:\*  
4: /cgn2\_6/prodata/2/lna/6B.COMB.seq:\*  
5: /cgn2\_6/prodata/2/lna/PCUTUS.COMB.seq:\*  
6: /cgn2\_6/prodata/2/lna/Backfiles1.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	26.8	2.7	30	4	US-09-358-972-102
2	26.8	2.7	30	4	US-09-358-972-103
3	26.8	2.7	30	4	US-09-406-147-32
4	26.8	2.7	30	4	US-09-406-147-34
5	20.8	2.1	24	1	US-08-587-209-6
6	20.8	2.1	24	1	US-08-689-236-6
7	20.8	2.1	24	1	US-08-689-236-6
8	20.8	2.1	24	1	US-08-692-725-6
9	20.8	2.1	24	2	US-08-692-726-6
10	20.8	2.1	58	2	US-08-431-527A-7
11	20.8	2.1	58	4	US-09-214-278-30
12	20.6	2.1	52	3	US-08-886-967-3
13	20.6	2.1	52	4	US-09-306-949-3
14	20.6	2.1	59	3	US-08-874-825-118
15	20.6	2.1	50	2	US-08-450-905B-7
16	20.6	2.1	50	3	US-07-982-759F-7
17	20.6	2.1	57	1	US-08-141-892A-4
18	20.6	2.1	57	2	US-08-583-447A-4
19	20.6	2.1	57	2	US-08-467-920-4
20	20.6	2.1	57	3	US-08-635-930-4
21	20.6	2.1	57	3	US-09-193-997-4
22	20.6	2.1	57	4	US-09-138-237A-4
23	20.6	2.1	60	1	US-08-478-370-4
24	20.6	2.1	43	1	US-07-959-946-12
25	20.6	2.1	43	1	US-08-333-577-12
26	20.6	2.1	43	5	PCT-US92-08634-12
27	20.6	2.1	58	1	US-08-105-483-174

C 28	19.6	2.0	58	1	US-08-709-209-174	Sequence 174, App
C 29	19.6	2.0	58	1	US-08-303-275-62	Sequence 62, App1
C 30	19.6	2.0	58	1	US-08-458-101-174	Sequence 174, App
C 31	19.6	2.0	60	1	US-07-670-296-19	Sequence 19, App1
C 32	19.6	2.0	60	1	US-08-093-781-20	Sequence 20, App1
C 33	19.6	2.0	60	3	US-08-963-602-2	Sequence 2, App1
C 34	19.4	1.9	37	2	US-08-403-853-8	Sequence 8, App1
C 35	19.4	1.9	60	1	US-08-487-890A-127	Sequence 127, App
C 36	19.4	1.9	60	2	US-08-478-435-127	Sequence 127, App
C 37	19.4	1.9	60	2	US-08-337-483-127	Sequence 127, App
C 38	19.4	1.9	60	2	US-08-478-373-127	Sequence 127, App
C 39	19.4	1.9	60	3	US-08-474-671-127	Sequence 127, App
C 40	19.4	1.9	60	3	US-08-483-577A-127	Sequence 127, App
C 41	19.4	1.9	60	4	US-08-897-438-127	Sequence 127, App
C 42	19.2	1.9	58	2	US-08-431-527A-6	Sequence 6, App1
C 43	19.2	1.9	60	1	US-07-609-716-72	Sequence 72, App1
C 44	19.2	1.9	60	3	US-08-475-411A-72	Sequence 72, App1
C 45	19.2	1.9	60	4	US-08-478-029A-72	Sequence 72, App1

## ALIGNMENTS

RESULT 1  
US-09-358-972-102/c  
; Sequence 102, Application US/09358972  
; Patent No. 6235480  
; GENERAL INFORMATION:  
; APPLICANT: Shultz, John W.  
; APPLICANT: Lewis, Martin K.  
; APPLICANT: Liepp, Donna  
; APPLICANT: Mandrekar, Michelle  
; APPLICANT: Kephart, Daniel  
; APPLICANT: Rhodes, Richard B.  
; APPLICANT: Andrews, Christine A.  
; APPLICANT: Hartnett, James R.  
; APPLICANT: Gu, Trent  
; APPLICANT: Olson, Ryan J.  
; APPLICANT: Wood, Keith W.  
; APPLICANT: Welch, Roy  
; TITLE OF INVENTION: Nucleic Acid Detection  
; FILE REFERENCE: PRO-103 6868/75528  
; CURRENT APPLICATION NUMBER: US/09/358,972  
; CURRENT FILING DATE: 1999-07-22  
; EARLIER APPLICATION NUMBER: 09/252,436  
; EARLIER FILING DATE: 1999-02-18  
; EARLIER APPLICATION NUMBER: 09/042,287  
; EARLIER FILING DATE: 1998-03-13  
; NUMBER OF SEQ ID NOS: 290  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 102  
; LENGTH: 30  
; TYPE: DNA  
; ORGANISM: Campylobacter jejuni  
; FEATURE:  
; OTHER INFORMATION: probe to Campylobacter jejuni  
US-09-358-972-102

Query Match 2.7%; Score 26.8; DB 4; Length 30;  
Best local similarity 93.3%; Pred. No. 82;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatgtcaaaagcttaaaacagact 318  
||||| ||||| ||||| ||||| |||||  
DB 30 CAAGATGACAAAGTTTAAAAACAAGACT 1

RESULT 2  
US-09-358-972-103  
; Sequence 103, Application US/09358972  
; Patent No. 6235480  
; GENERAL INFORMATION:

APPLICANT: Shultz, John W  
APPLICANT: Lewis, Martin K.  
APPLICANT: Lieppe, Donna  
APPLICANT: Mandrekar, Michelle  
APPLICANT: Kephart, Daniel  
APPLICANT: Rhodes, Richard B.  
APPLICANT: Andrews, Christine A.  
APPLICANT: Hartnett, James R.  
APPLICANT: Gu, Trent  
APPLICANT: Olson, Ryan J.  
APPLICANT: Wood, Keith W.  
APPLICANT: Welch, Roy  
TITLE OF INVENTION: Nucleic Acid Detection  
FILE REFERENCE: Pro-103 6868/75528  
CURRENT APPLICATION NUMBER: US/09/358,972  
CURRENT FILING DATE: 1999-07-22  
EARLIER APPLICATION NUMBER: 09/252,436  
EARLIER FILING DATE: 1999-02-18  
EARLIER APPLICATION NUMBER: 09/042,287  
EARLIER FILING DATE: 1998-03-13  
NUMBER OF SEQ ID NOS: 290  
SOFTWARE: Patentln Ver. 2.0  
SEQ ID NO 103  
LENGTH: 30  
TYPE: DNA  
ORGANISM: Campylobacter jejuni  
FEATURE:  
OTHER INFORMATION: probe to Campylobacter jejuni  
US-09-358-972-103

Query Match 2.7%: Score 26.8; DB 4; Length 30;  
Best Local Similarity 93.3%; Pred. No. 82;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 289 caagatgtcgaagcttaaaacaagaact 318  
||||| ||||| ||||| ||||| |||||  
Db 1 caagatgacacaagtttaaaacaagaact 30

RESULT 3  
US-09-406-147-32/c  
Sequence 32, Application US/09406147  
Patent No. 6270974  
GENERAL INFORMATION:  
APPLICANT: Shultz, John W  
APPLICANT: Lewis, Martin K  
APPLICANT: Lieppe, Donna  
APPLICANT: Mandrekar, Michelle  
APPLICANT: Kephart, Daniel  
APPLICANT: Rhodes, Richard B  
APPLICANT: Andrews, Christine A  
APPLICANT: Hartnett, James R  
APPLICANT: Gu, Trent  
APPLICANT: Wood, Keith V  
APPLICANT: Welch, Roy  
TITLE OF INVENTION: EXOGENOUS NUCLEIC ACID DETECTION  
FILE REFERENCE: EXOGENOUS NUCLEIC ACID DETECTION  
CURRENT APPLICATION NUMBER: US/09/406,147  
CURRENT FILING DATE: 1999-09-27  
EARLIER APPLICATION NUMBER: 09/252,436  
EARLIER FILING DATE: 1999-02-18  
EARLIER APPLICATION NUMBER: 09/042,287  
EARLIER FILING DATE: 1998-03-13  
NUMBER OF SEQ ID NOS: 92  
SOFTWARE: Patentln Ver. 2.0  
SEQ ID NO 32  
LENGTH: 30  
TYPE: DNA  
ORGANISM: Campylobacter jejuni  
US-09-406-147-32

Query Match 2.7%: Score 26.8; DB 4; Length 30;  
Best Local Similarity 93.3%; Pred. No. 82;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 289 caagatgtcgaagcttaaaacaagaact 318  
||||| ||||| ||||| ||||| |||||  
Db 30 CAAGATGACAAAGTTTAAACAGAAGACT 1

RESULT 4  
US-09-406-147-34  
Sequence 34, Application US/09406147  
Patent No. 6270974  
GENERAL INFORMATION:  
APPLICANT: Shultz, John W  
APPLICANT: Lewis, Martin K  
APPLICANT: Lieppe, Donna  
APPLICANT: Mandrekar, Michelle  
APPLICANT: Kephart, Daniel  
APPLICANT: Rhodes, Richard B  
APPLICANT: Andrews, Christine A  
APPLICANT: Hartnett, James R  
APPLICANT: Gu, Trent  
APPLICANT: Wood, Keith V  
TITLE OF INVENTION: EXOGENOUS NUCLEIC ACID DETECTION  
FILE REFERENCE: EXOGENOUS NUCLEIC ACID DETECTION  
CURRENT APPLICATION NUMBER: US/09/406,147  
CURRENT FILING DATE: 1999-09-27  
EARLIER APPLICATION NUMBER: 09/252,436  
EARLIER FILING DATE: 1999-02-18  
EARLIER APPLICATION NUMBER: 09/042,287  
EARLIER FILING DATE: 1998-03-13  
NUMBER OF SEQ ID NOS: 92  
SOFTWARE: Patentln Ver. 2.0  
SEQ ID NO 34  
LENGTH: 30  
TYPE: DNA  
ORGANISM: Campylobacter jejuni  
US-09-406-147-34

Query Match 2.7%: Score 26.8; DB 4; Length 30;  
Best Local Similarity 93.3%; Pred. No. 82;  
Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 289 caagatgtcgaagcttaaaacaagaact 318  
||||| ||||| ||||| ||||| |||||  
Db 1 caagatgacacaagtttaaaacaagaact 30

RESULT 5  
US-08-587-209-6/c  
Sequence 6, Application US/08587209  
Patent No. 5612473  
GENERAL INFORMATION:  
APPLICANT: Wu, Linxian  
APPLICANT: Coombs, Jana  
APPLICANT: Malmstrom, Sharon L.  
APPLICANT: Glass, Michael J.  
TITLE OF INVENTION: Methods and Apparatus for Preparing, Amplifying,  
NUMBER OF SEQUENCES: 30  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David O. Seeley, Esq.  
ADDRESSEE: Workman, Nydegger & Seeley  
STREET: 1000 Eagle Gate Tower  
STREET: 60 East South Temple  
CITY: Salt Lake City  
STATE: Utah  
COUNTRY: USA  
ZIP: 84111  
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette, 3.50 inch,  
MEDIUM TYPE: 1.44 Mb storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 6.0a for WINDOWS  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/587,209  
FILING DATE: 16-JAN-1996  
CLASSIFICATION: 435  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 24 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-587-209-6

Query Match 2.1%; Score 20.8; DB 1; Length 24;  
Best Local Similarity 91.7%; Pred. No. 3e+03;  
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 91 ggccttagaactcaactccgcagca 114  
|||||  
DB 24 GGCTTAGAATTAACTCAGCAGCA 1

RESULT 6  
US-08-689-236-6/c  
Sequence 6, Application US/08689236  
Patent No. 5738995  
GENERAL INFORMATION:  
APPLICANT: Wu, Linxian  
APPLICANT: Coombs, Jana  
APPLICANT: Malmstrom, Sharon L.  
APPLICANT: Glass, Michael J.  
TITLE OF INVENTION: Methods and Apparatus for  
PREPARING, AMPLIFYING, AND DISCRIMINATING MULTIPLE ANALYTES  
NUMBER OF SEQUENCES: 30  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David O. Seeley, Esq.  
ADDRESSEE: Workman, Nydegger & Seeley  
STREET: 1000 Eagle Gate Tower  
STREET: 60 East South Temple  
CITY: Salt Lake City  
STATE: Utah  
COUNTRY: USA  
ZIP: 84111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.50 inch,  
MEDIUM TYPE: 1.44 Mb storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 6.0a for WINDOWS  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/689,236  
FILING DATE: 16-JAN-1996  
CLASSIFICATION: 435  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 24 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-689-236-6

Query Match 2.1%; Score 20.8; DB 1; Length 24;  
Best Local Similarity 91.7%; Pred. No. 3e+03;  
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 91 ggccttagaactcaactccgcagca 114  
|||||  
DB 24 GGCTTAGAATTAACTCAGCAGCA 1

RESULT 7  
US-08-689-235-6/c  
Sequence 6, Application US/08689235  
Patent No. 5753444  
GENERAL INFORMATION:  
APPLICANT: Wu, Linxian  
APPLICANT: Coombs, Jana  
APPLICANT: Malmstrom, Sharon L.  
APPLICANT: Glass, Michael J.  
TITLE OF INVENTION: Methods and Apparatus for  
PREPARING, AMPLIFYING, AND DISCRIMINATING MULTIPLE ANALYTES  
NUMBER OF SEQUENCES: 30  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David O. Seeley, Esq.  
ADDRESSEE: Workman, Nydegger & Seeley  
STREET: 1000 Eagle Gate Tower  
STREET: 60 East South Temple  
CITY: Salt Lake City  
STATE: Utah  
COUNTRY: USA  
ZIP: 84111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.50 inch,  
MEDIUM TYPE: 1.44 Mb storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 6.0a for WINDOWS  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/689,235  
FILING DATE: 16-JAN-1996  
CLASSIFICATION: 435  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 24 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-689-235-6

Query Match 2.1%; Score 20.8; DB 1; Length 24;  
Best Local Similarity 91.7%; Pred. No. 3e+03;  
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 91 ggccttagaactcaactccgcagca 114  
|||||  
DB 24 GGCTTAGAATTAACTCAGCAGCA 1

RESULT 8  
US-08-692-725-6/c  
Sequence 6, Application US/08692725  
Patent No. 5756701  
GENERAL INFORMATION:  
APPLICANT: Wu, Linxian  
APPLICANT: Coombs, Jana  
APPLICANT: Malmstrom, Sharon L.  
APPLICANT: Glass, Michael J.  
TITLE OF INVENTION: Methods and Apparatus for  
PREPARING, AMPLIFYING, AND DISCRIMINATING MULTIPLE ANALYTES  
NUMBER OF SEQUENCES: 30  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David O. Seeley, Esq.  
ADDRESSEE: Workman, Nydegger & Seeley  
STREET: 1000 Eagle Gate Tower  
STREET: 60 East South Temple  
CITY: Salt Lake City

```

; STANDEDNESS: 31
; TOPOLOGY: linear
; MOLECULE TYPE: DNA

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US-08-431-527A-7

APPLICANT: Sakano, Seiji

```

RESULT 11
US-09-214-278-30/c
; Sequence 30, Application US/09214278
; Patent No. 6291210
; GENERAL INFORMATION:
; APPLICANT: Sakano, Sei-ji
; APPLICANT: Ito, Akira

```



;; TITLE OF INVENTION: DIFFERENTIATION-SUPPRESSIVE POLYPEPTIDE  
;; FILE REFERENCE: KP-8576  
;; CURRENT APPLICATION NUMBER: US/09/214,278  
;; NUMBER OF SEQ ID NOS: 32  
;; SOFTWARE: PatentIn Ver. 2.1  
;; SEQ ID NO 30  
;; LENGTH: 58  
;; TYPE: DNA  
;; ORGANISM: Artificial Sequence  
;; OTHER INFORMATION: Description of Artificial Sequence: synthetic DNA  
US-09-214-278-30

Query Match 2.1%; Score 20.8; DB 4; Length 58;  
Best Local Similarity 78.1%; Pred. No. 4.3e+03;  
Matches 25; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Oy 769 gtaaggtcgtatcatcagatggtcagagaa 800  
|| ||| ||||| ||||| ||||| ||  
Db 42 GGCAGAGAGATTTAAAGATGATGATGATTA 11

RESULT 12  
US-08-886-967-3/c  
; Sequence 3, Application US/08886967  
; Patent No. 6068993  
; GENERAL INFORMATION:  
; APPLICANT: ASTOLFI, SPARTACO  
; APPLICANT: DE LIMA, BEATRIZ D.  
; APPLICANT: THIEMANN, JOSEF E.  
; APPLICANT: TUNES DE SOUSA, HELOISA R.  
; APPLICANT: VILELA, LUCIANO  
; TITLE OF INVENTION: VECTOR FOR EXPRESSION OF HETEROLOGOUS  
; TITLE OF INVENTION: PROTEIN AND METHODS FOR EXTRACTING RECOMBINANT PROTEIN AND  
; TITLE OF INVENTION: FOR PURIFYING ISOLATED RECOMBINANT INSULIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: FROMMER LAWRENCE & HAUG LLP  
; STREET: 745 FIFTH AVENUE  
; CITY: NEW YORK  
; STATE: NEW YORK  
; COUNTRY: USA  
; ZIP: 10151  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/886,967  
; FILING DATE: 02-JUL-1997  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: HAUG, EDGAR H.  
; REGISTRATION/DOCKET NUMBER: 29,309  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 212-588-0800  
; TELEFAX: 212-588-0500  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 52 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-886-967-3

Query Match 2.1%; Score 20.6; DB 3; Length 52;  
Best Local Similarity 67.4%; Pred. No. 4.6e+03;

Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;  
Oy 400 ttaagtgtggtttaccatcagaattccaatcggttaa 442  
||||| ||| ||| ||||| ||| ||| ||  
Db 52 TTAAGTGTACTTAATAGAGAGAAATTTCATTGTTTAA 10

RESULT 13  
US-09-306-949-3/c  
; Sequence 3, Application US/09306949  
; Patent No. 6281329  
; GENERAL INFORMATION:  
; APPLICANT: ASTOLFI, SPARTACO  
; APPLICANT: DE LIMA, BEATRIZ D.  
; APPLICANT: THIEMANN, JOSEF E.  
; APPLICANT: TUNES DE SOUSA, HELOISA R.  
; APPLICANT: VILELA, LUCIANO  
; TITLE OF INVENTION: VECTOR FOR EXPRESSION OF HETEROLOGOUS  
; TITLE OF INVENTION: PROTEIN AND METHODS FOR EXTRACTING RECOMBINANT PROTEIN AND  
; TITLE OF INVENTION: FOR PURIFYING ISOLATED RECOMBINANT INSULIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: FROMMER LAWRENCE & HAUG LLP  
; STREET: 745 FIFTH AVENUE  
; CITY: NEW YORK  
; STATE: NEW YORK  
; COUNTRY: USA  
; ZIP: 10151  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/306,949  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/886,967  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: HAUG, EDGAR H.  
; REGISTRATION/DOCKET NUMBER: 29,309  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 212-588-0800  
; TELEFAX: 212-588-0500  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 52 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-09-306-949-3

Query Match 2.1%; Score 20.6; DB 4; Length 52;  
Best Local Similarity 67.4%; Pred. No. 4.6e+03;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

Oy 400 ttaagtgtggtttaccatcagaattccaatcggttaa 442  
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Db 52 TTAAGTGTACTTAATAGAGAGAAATTTCATTGTTTAA 10

RESULT 14  
US-08-874-825-118/c  
; Sequence 118, Application US/08874825  
; Patent No. 6057101  
; GENERAL INFORMATION:  
; APPLICANT: Nandabalan, Krishnan  
; APPLICANT: Rothberg, Jonathan

APPLICANT: Yang, Melja  
APPLICANT: Knight, James  
TITLE OF INVENTION: IDENTIFICATION AND COMPARISON OF  
TITLE OF INVENTION: PROTEIN-PROTEIN INTERACTIONS THAT OCCUR IN POPULATIONS  
TITLE OF INVENTION: AND IDENTIFICATION OF INHIBITORS OF THESE INTERACTIONS  
NUMBER OF SEQUENCES: 122  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Pennie & Edmonds  
STREET: 1155 Avenue of the Americas  
CITY: New York  
STATE: NY  
COUNTRY: USA  
ZIP: 10036/2711  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/874,825  
FILING DATE: 13-JUN-1997  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/663,824  
FILING DATE: 14-JUN-1996  
ATTORNEY/AGENT INFORMATION:  
NAME: Mirock, S. Leslie  
REGISTRATION NUMBER: 18,872  
REFERENCE/DOCKET NUMBER: 7934-045  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 212-790-9090  
TELEFAX: 212-869-8864  
TELEX: 66141 PENNIE  
INFORMATION FOR SEQ ID NO: 118:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 39 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
US-08-874-825-118

Query Match 2.0%; Score 20; DB 3; Length 39;  
Best Local Similarity 82.1%; Pred. No. 6e+03;  
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 768 aggtgaagtgatcattcagatggtgat 795  
|||||  
DB 39 AGGTCAGGTCCTTCACATGCTCAT 12

RESULT 15  
US-08-450-905B-7/c  
Sequence 7, Application US/08450905B  
Patent No. 5856301  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: Stem Cell Inhibiting Proteins  
NUMBER OF SEQUENCES: 178  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HALE and DORR  
STREET: 60 State Street  
CITY: Boston  
STATE: MA  
ZIP: 02109  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/450,905B  
FILING DATE: 26-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/982,759  
FILING DATE: 08-MAR-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: GB 9127319.3  
FILING DATE: 23-DEC-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: GB 9221587.0  
FILING DATE: 14-OCT-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: BAKER, HOLIE L.  
REGISTRATION NUMBER: 31,321  
REFERENCE/DOCKET NUMBER: 102,378.120DV-2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617-526-6110  
TELEFAX: 617-526-5000  
INFORMATION FOR SEQ ID NO: 7:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 50 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
FEATURE:  
NAME/KEY: misc.feature  
LOCATION: 1..50  
OTHER INFORMATION: /product- "OLIGOMER FOR  
OTHER INFORMATION: CONSTRUCTION OF SYNTHETIC LD78 GENE"  
US-08-450-905B-7

Query Match 2.0%; Score 20; DB 2; Length 50;  
Best Local Similarity 72.2%; Pred. No. 6.6e+03;  
Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

OY 368 caaatcaccacttcaatggaacaacttta 403  
|||||  
DB 36 CAAATTCACAAATTCATGCTGACTACTTGAA 1

Search completed: April 17, 2002, 02:18:56  
Job time: 3297 sec

wed Apr 17 07:36:45 2002

us-09-439-311-1.rtf

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GenCore version 4.5  
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 17, 2002, 01:16:14 : Search time 1464.96 Seconds  
(without alignments)  
7327.868 Million cell updates/sec

Title: US-09-439-311-1

Perfect score: 999

Sequence: 1 attacacacaaatgttcgacg.....ttaaaatgctgtagagat 999

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 11351937 seqs, 5372889281 residues

Total number of hits satisfying chosen parameters: 111874

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Maximum Match 0%

Listing first 45 summaries

Database :

EST:\*  
1: em\_estfun:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estom:\*  
5: em\_estpl:\*  
6: em\_estba:\*  
7: em\_estro:\*  
8: em\_estov:\*  
9: em\_hic:\*  
10: gp\_estl:\*  
11: gp\_estl2:\*  
12: gp\_hic:\*  
13: gp\_gss:\*  
14: em\_gss\_fun:\*  
15: em\_gss\_hum:\*  
16: em\_gss\_hiv:\*  
17: em\_gss\_pln:\*  
18: em\_gss\_pro:\*  
19: em\_gss\_rod:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_other:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	23.6	2.2	50	10	AU104183	AU104183 AU104183
2	21.6	2.4	60	10	AA608932	AA608932 af03d01.s
3	21.21	2.1	52	10	AA122474	AA122474 mg27h10.r
4	20.8	2.1	59	10	AM687832	AM687832 NF01A01R
5	20.8	2.1	60	11	BF638864	BF638864 NF06H02P
6	20.6	2.1	58	10	AA874675	AA874675 vW84C07.r
7	20.4	2.0	51	13	AZ812517	AZ812517 2M0079C15
8	20.4	2.0	55	10	AU104315	AU104315 AU104315
9	20.2	2.0	50	10	AU103731	AU103731 AU103731
10	20.2	2.0	54	13	AZ954544	AZ954544 2M0220G15
11	20.2	2.0	58	11	BF131272	BF131272 601819529
12	20.2	2.0	59	11	W38842	W38842 zb28c08.r1

13	20	2.0	54	13	AZ434413	AZ434413 IM0220118
14	20	2.0	56	10	AM780772	AM780772 s185c06.y
15	20	2.0	56	13	AF149647	AF149647 AF149647.y
16	20	2.0	56	13	AZ451302	AZ451302 IM0250012
17	20	2.0	58	10	AA164130	AA164130 mq84f08.r
18	20	2.0	60	10	AA878830	AA878830 of83f08.s
19	20	2.0	60	13	AZ778864	AZ778864 2M0014P24
20	20	2.0	60	13	B44939	B44939 HS-1060-A2-
21	19.8	2.0	50	11	BC939021	BC939021 cn30a04.y
22	19.8	2.0	58	11	BC058837	BC058837 nag43c06.
23	19.8	2.0	60	10	AU060311	AU060311 AU060311
24	19.6	2.0	50	13	AZ397298	AZ397298 IM0162A13
25	19.6	2.0	54	10	AM307272	AM307272 sf54h07.y
26	19.6	2.0	59	13	AZ626633	AZ626633 IM0467C04
27	19.4	1.9	50	10	AU107058	AU107058 AU107058
28	19.4	1.9	52	10	AM687820	AM687820 NF013G12R
29	19.4	1.9	53	11	BG370297	BG370297 na129h10.
30	19.4	1.9	56	13	AM513614	AM513614 xo47d06.x
31	19.4	1.9	56	13	AZ608724	AZ608724 IM0433C09
32	19.4	1.9	58	10	AA190288	AA190288 mt93g05.r
33	19.4	1.9	58	11	T18527	T18527 hbc2089 Hum
34	19.4	1.9	58	11	T18565	T18565 hbc2087 Hum
35	19.4	1.9	59	10	A1855540	A1855540 sc20e08.Y
36	19.2	1.9	51	11	D18206	D18206 MUGS00476
37	19.2	1.9	52	10	AA492773	AA492773 v176h09.r
38	19.2	1.9	54	13	AZ783521	AZ783521 2M0025P17
39	19	1.9	50	11	BC271613	BC271613 na159h03.
40	19	1.9	52	10	AM396361	AM396361 sh27b06.y
41	19	1.9	53	11	BG408927	BG408927 g8b3c07.y
42	19	1.9	54	13	AZ345736	AZ345736 HUMGS000792
43	19	1.9	60	11	C01215	C01215 HUMGS000792
44	19	1.9	60	11	BF638586	BF638586 NE054G03P
45	18.8	1.9	50	10	AU104948	AU104948 AU104948

## ALIGNMENTS

RESULT 1  
AU104183/c  
LOCUS  
DEFINITION  
AU104183 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone  
KAT06093, mRNA sequence.  
ACCESSION  
AU104183  
VERSION  
AU104183.1 GI:13553704  
KEYWORDS  
EST.  
SOURCE  
human.  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE  
1 (bases 1 to 50)  
Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata  
,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo  
,K., Suyama,A. and Sugano,S.  
Fine Structural analysis of transcription start sites of human  
mRNAs using full-length enriched and 5'-end enriched cDNA libraries  
unpublished (2001)  
JOURNAL  
Contact: Yutaka Suzuki  
Department of Virology  
Institute of Medical Science, University of Tokyo  
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan  
Email: yusuzuki@ims.u-tokyo.ac.jp  
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano  
,S. Construction and characterization of a full length-enriched and  
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).  
FEATURES  
source  
1. 50  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="KAT06093"  
/clone\_lib="Sugano Homo sapiens cDNA library"  
BASE COUNT  
12 a 12 c 4 g 22 t  
ORIGIN

Query Match 2.4%; Score 23.6; DB 10; Length 50;  
 Best Local Similarity 69.6%; Pred. No. 1.5e+05;  
 Matches 32; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

Oy 902 gagggatataatcacaggtagcatagtgtagagctgtatatt 947  
 Db 46 GAGGGTTTAAATTAAGTTAGACTGGAATTAAGCTACATGCTT 1

RESULT 2  
 AA608932 60 bp mRNA EST 02-MAR-1998  
 LOCUS af03001.s1 Soares\_testis\_NHT Homo sapiens cDNA clone IMAGE:1030561  
 DEFINITION 3, similar to gb:M10058 ASIALOGLYCOPROTEIN RECEPTOR 1 (HUMAN),  
 mRNA sequence.  
 ACCESSION AA608932  
 VERSION AA608932.1 GI:2457360  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 REFERENCE 1 (bases 1 to 60)  
 AUTHORS Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisels,G., Jost,S.,  
 Krizman,D., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M., Martin,  
 J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F., Theising,B.,  
 White,Y., Wylie,T., Waterston,R. and Wilson,R.  
 TITLE Washu-NCI human EST Project  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Wilson RK  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: estewatson.wustl.edu  
 This clone is available royalty-free through LNL; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
 Trace considered overall poor quality  
 Insert length: 1146 Std Error: 0.00  
 Seq primer: -40m13 fwd. ET from Amersham  
 High quality sequence stop: 1.  
 Location/Qualifiers  
 1. 60  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone\_id="IMAG:1030561"  
 /clone\_lib="Soares\_testis\_NHT"  
 /sex="male"  
 /lab\_host="DH10B"  
 /note="Vector: PT73D-Pac (Pharmacia) with a modified  
 polylinker. Site 1: Not I. Site 2: Eco RI. 1st strand cDNA  
 was prepared from mRNA obtained from Clontech Laboratories  
 , Inc., and primed with a Not I - oligo(dT) primer [5'  
 TGTACCAATCTGAGTGGAGCGGCCCAATTTTCTTTTCTTTT 3']  
 Double-stranded cDNA was ligated to Eco RI adaptors  
 (Pharmacia), digested with Not I and cloned into the Not I  
 and Eco RI sites of the modified pT73 vector. Library  
 went through one round of normalization to Cot5, and was  
 constructed by Bento Soares and M. Fatima Bonalido."

BASE COUNT 16 a 20 c 11 g 13 t

ORIGIN

Query Match 2.2%; Score 21.6; DB 10; Length 60;  
 Best Local Similarity 63.5%; Pred. No. 4.5e+05;  
 Matches 33; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

Oy 205 ggtatcttgcaactgcagataagcctatgcatgacacttaaaccttag 256  
 Db 9 GGTAACTGCACTGCAAGAAACGCCAGCGGTTTCAAGCTCTCACACTTCG 60

RESULT 3  
 AA122474 52 bp mRNA EST 13-FEB-1997  
 LOCUS mg27h10.r1 Barstead MRLB1 Mus musculus cDNA clone IMAGE:580003 5'  
 DEFINITION similar to SW:COX3\_MOUSE P00416 CYTOCHROME C OXIDASE POLYPEPTIDE  
 ITT, mRNA sequence.  
 ACCESSION AA122474  
 VERSION AA122474.1 GI:1681537  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus  
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE 1 (bases 1 to 52)  
 AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,  
 Geisels,G., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,  
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,  
 Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and  
 Waterston,R.  
 TITLE The Washu-HMI Mouse EST Project  
 JOURNAL Unpublished (1996)  
 COMMENT Contact: Marra M/Mouse EST Project  
 Washu-HMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: mouseest@watson.wustl.edu  
 This clone is available royalty-free through LNL; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
 MGI:354651  
 Trace considered overall poor quality  
 Possible reversed clone: similarity on wrong strand  
 Seq primer: -28m13 rev2 ET from Amersham  
 High quality sequence stop: 1.  
 Location/Qualifiers  
 1. 52  
 /organism="Mus musculus"  
 /strain="BALB/c"  
 /db\_xref="taxon:10090"  
 /clone\_id="IMAG:580003"  
 /clone\_lib="Barstead MRLB1"  
 /sex="mixed"  
 /tissue\_type="kidney"  
 /dev\_stage="6 weeks"  
 /lab\_host="DH10B"  
 /note="Vector: PT73D-Pac (Pharmacia) with a modified  
 polylinker. Site 1: EcoRI. Site 2: NotI. 1st strand cDNA  
 was primed with a Not I - oligo(dT) primer [5'  
 TGTACGAATCTGAGTGGAGCGGCCCTTTTCTTTTCTTTTCTTTT 3']  
 double-stranded cDNA was ligated to Eco RI adaptors  
 (CATGATTCGGTACC), digested with Not I and cloned into the  
 Not I and Eco RI sites of the modified pT73 vector.  
 Library constructed by Bob Barstead."

BASE COUNT 20 a 7 c 8 g 17 t

ORIGIN

Query Match 2.1%; Score 21; DB 10; Length 52;  
 Best Local Similarity 73.0%; Pred. No. 6.2e+05;  
 Matches 27; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Oy 810 aatttcagctacatgctgtaaaagatacaactcgt 846  
 Db 4 AATTTCACAATTATATCTCTAGAGAGTTGCAATAGTT 40

RESULT 4  
 AM687832 59 bp mRNA EST 20-DEC-2000  
 LOCUS NF014A01RTJF1003 Developing root Medicago truncatula cDNA clone  
 DEFINITION NF014A01RT 5', mRNA sequence.

REFERENCE	AM687832	GI:11929203
VERSION	AM687832.2	GI:11929203
KEYWORDS	EST.	
SOURCE	barrel medic.	
ORGANISM	Medicago truncatula	
REFERENCE	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifoliaceae; Medicago.	
AUTHORS	1 (bases 1 to 59)	
TITLE	Watson,B.S., Shih,H.-S., Lopez-Meyer,M., Scott,A.D., Harris,A.R., Gonzales,R.A., Bell,C.J., Flores,H.R., Imman,J.T., Weller,J.W., May,G.D. and Palva,N.L.	
JOURNAL	Expressed Sequence Tags from the Samuel Roberts Noble Foundation Medicago truncatula root library	
COMMENT	Unpublished (2000) On Apr 14, 2000 this sequence version replaced gi:7562568. Contact: Palva NL Plant Biology Division The Samuel Roberts Noble Foundation 2510 Sam Noble Parkway, Ardmore, OK 73402, USA Tel: 580 221 7317 Fax: 580 221 7380 Email: nlpalva@noble.org Insert Length: 721 std Error: 0.00 Plate: 014 row: A column: 01 Seq primer: TCACACGACGAAACGCTATGAC. Location/Qualifiers 1..59	
FEATURES	/organism="Medicago truncatula" /db_xref="taxon:3880" /clone="NF014A01R" /clone_11b="Developing root" /tissue_type="root" /dev_stage="Pooled developmental" /note="Vector: Lambda zap; Total RNA was extracted from non-nodulated roots of plants grown in 1 mM nitrate medium. Samples were taken at four time points (approximately two days, one, two and six weeks post germination) representing early seedling growth to nitrogen limitation."	
BASE COUNT	19 a 13 g 18 t	
ORIGIN		
Query Match	2.1%; Score 20.8; DB 10; Length 59;	
Best Local Similarity	70.0%; Pred. No. 6.9e+05;	
Matches	28; Conservative 0; Mismatches 12; Indels 0; Gaps 0;	
QY	822 caatgcgtctaaagatacaactggtcttaacgctcttaaa 861	
Db	54 CATTGCATATGACGAAACACACAGCTGTCATATCACTTCA 15	
RESULT		
LOCUS	BF638864 60 bp mRNA EST 19-DEC-2000	
DEFINITION	NF060H02PL1F1026 phosphate starved leaf Medicago truncatula cDNA	
ACCESSION	clone NF060H02PL 5', mRNA sequence.	
VERSION	BF638864	
KEYWORDS	BF638864.1 GI:11903022	
ORGANISM	EST.	
SOURCE	barrel medic.	
REFERENCE	Medicago truncatula	
AUTHORS	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifoliaceae; Medicago.	
TITLE	1 (bases 1 to 60)	
COMMENT	Lin,J., Scott,A.D., Harris,A.R., Gonzales,R.A., Bell,C.J., Flores,H.R., Imman,J.T., Weller,J.W., May,G.D. and Harrison,M.J. Expressed Sequence Tags from the Samuel Roberts Noble Foundation Medicago truncatula phosphate-starved leaf library	

JOURNAL COMMENT

Unpublished (2000)  
Contact: Harrison MJ  
Plant Biology Division  
The Samuel Roberts Noble Foundation  
2510 Sam Noble Parkway, Ardmore, OK 73402, USA  
Tel: 580 221 7325  
Fax: 580 221 7380  
Email: mjharrison@noble.org  
Insert Length: 60 Std Error: 0.00  
Plate: 060 row: H column: 02  
Seq primer: TCACACAGCAAAACGCTATGC.

FEATURES

Source location/qualifiers

1..60  
/organism="Medicago truncatula"  
/db\_xref="taxon:3880"  
/clone="NF060H02PL"  
/clone\_lib="Phosphate starved leaf"  
/tissue\_type="leaf"  
/dex\_stage="trifoliolate"  
/note="Vector: Lambda Zap; At the trifoliolate stage, M. truncatula plants were transplanted to phosphate-free sand and grown for a further 30 days. During this 30 day period, the plants were fertilized twice weekly with 1/2 Hoaglands solution containing only 20mM potassium phosphate. RNA was prepared from above ground tissues."

BASE COUNT

15 a 17 c 12 g 9 t 7 others

ORIGIN

Query Match 2.1%; Score 20.8; DB 11; Length 60;  
Best Local Similarity 59.6%; Pred. No. 6.9e+05;  
Matches 28; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

Qy 76 tcaaacctcagcttcaggcttagaatcaactccgacgatcatgc 122  
||| ||| | | | | | | | | | | | | | | |  
Db 47 tcagacctnngtttnaggttgattgatgcncagctnmacngmtcgctggtc 1

RESULT 6  
AA874675/c

LOCUS  
DEFINITION  
IMAGE:1261644 5' similar to TR:003713 003713 CYTOCHROME B ;, mRNA

ACCSSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
house mouse.  
Mus musculus  
Eukaryota; Metazoa; Chordata; Vertebrate; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 58)  
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,  
Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,  
Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,  
Thaising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and  
Waterston, R.  
The Mashu-HMMI Mouse EST Project  
Unpublished (1996)  
Contact: Marra M/Mouse EST Project  
WashU-HMMI Mouse EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: mouseest@wustl.edu  
This clone is available royalty-free through LINL; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
MGI:664196  
Trace considered overall poor quality  
Possible reversed clone: similarity on wrong strand  
Seq primer: -26m13 rev1 ET from Amersham  
High quality sequence stop: 1.

```

FEATURES
source
location/Qualifiers
1. 58
/organism="Mus musculus"
/strain="C57BL/6"
/db_xref="taxon:10090"
/clone_image:1261644"
/clone_1lb="Stratagene mouse skin (#937313)"
/sex="females"
/tissue_type="whole skin"
/dev_strage="11 weeks old"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: skin; Vector: pBluescript SK-; Site: 1; EcoRI
dt. Site: 2; XhoI; Cloned unidirectionally. Primer: Oligo
Whole skin from 11 week old C57BL/6 female mice.
Average insert size: 1.0 kb; Uni-ZAP XR Vector: -5'
adaptor sequence: 5' GAATTCGGCAGCAG 3' -3' adaptor
sequence: 5' CTCGAGTTTTTTTTTTTTTTT 3'"
19 a 18 c 2 g 19 t

```

[illegible]

musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMDA2 (g14732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

Query Match 2.0%: Score 20.4; DB 13; Length 51;  
Best Local Similarity 71.1%: Pred. No. 8.6e+05;  
Matches 27; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

Db 127 99gatgcagatagcagatagtttaagatctcaagcaca 164  
||||| ||| ||||| ||||| ||||| ||  
50 GGGAGCCCCGAGGCGAAGATTGTTGAGCTCAGGCGCA 13

RESULT 8  
LOCUS AU014315 mRNA EST 03-AUG-1998  
DEFINITION AU014315 Schizosaccharomyces pombe late log phase cDNA  
Schizosaccharomyces pombe cDNA clone spc09537, mRNA sequence.  
ACCESSION AU014315  
VERSION AU014315.1 GI:3369106  
KEYWORDS EST.  
SOURCE fission yeast.  
ORGANISM Schizosaccharomyces pombe  
Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;  
Schizosaccharomycetales; Schizosaccharomycetaceae;  
Schizosaccharomyces.  
1 (bases 1 to 55)  
Moriyomo,M. and Mita,K.  
Identification of expressed sequence tags of Schizosaccharomyces  
pombe  
Unpublished (1998)  
Contact: Mitsuoki Moriyomo  
Genome Research Group  
National Institute of Radiological Sciences  
9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan  
Email: moriyomo@nirs.go.jp

FEATURES  
Source Location/Qualifiers  
1..55  
/organism="Schizosaccharomyces pombe"  
/strain="972"  
/db\_xref="taxon:4896"  
/clone="spc09537"  
/clone\_lib="Schizosaccharomyces pombe late log phase cDNA"  
/sex="h minus"  
/note="Vector: M13mp19; The cDNA library of  
Schizosaccharomyces pombe was prepared by cloning cDNA  
into the SmaI site of M13mp19 DNA and the direction of DNA  
sequences was not always from 5' to 3'. The cDNA data of  
Schizosaccharomyces pombe are available for searching on  
the World Wide Web. (URL, <http://www.nirs.go.jp>)"

BASE COUNT 24 a 9 g 20 t  
ORIGIN  
Query Match 2.0%: Score 20.4; DB 10; Length 55;







```

/db_xref="taxon:10090"
/clone="UMGC1M0220118"
/clone_lib="Mouse 10kb plasmid UMGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/notes="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1147321141gb1Arl29072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

```

Query Match	2.0%	Score 20	DB 13	Length 54
Best Local Similarity	61.5%	Pred. No. 1	1e+06	
Matches	32	Conservative	0	Mismatches 20
				Indels 0
				Gaps 0
Qy	836	atacaactcgtgttc	caagcctctcaagaatgaaatg	gtgtaacctgttccttac 887
Db	2	ATTCAAAATGCTTTATGTC	TCTCAAAAATGTTAAAGATCAAT	GTGTCCTCAAC 53

AM780772	AM780772	56 bp	mRNA	EST	12-MAY-2000
LOCUS					
DEFINITION	g185cc06.y1 Gm-cl037 Glycine max CDNA clone			GENOME SYSTEMS CLONE	ID
ACCESSION	AM780772				
VERSION	AM780772.1	GI:7795447			
KEYWORDS	EST.				
ORGANISM	soybean.				
SOURCE	Glycine max				

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 56)	Shoemaker, R., Kelm, P., Vockin, L., Eppelding, J., Coryell, V., Khanna 'A., Bolla, B., Marrs, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wyle, T., Underwood, K., Steptoe, M., Thelsting, B., Allen, M., Bowers 'Y., Pearson, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk 'R., Ritter, E., Kohn, S., Shiu, T., Jackson, Y., Cardenas, M., McCann 'R., Waterson, R. and Wilson, R.	Public Soybean EST Project	Unpublished (1999)	Contact: Shoemaker R/Public Soybean EST Project

Small: est@watson.wustl.edu  
This clone is available through: Genome Systems, Inc. 4633 World Parkway Circle St. Louis, Missouri 63134 For further information call: (800) 430-0030 or (314) 427-3222 FAX: (888) 919-3324 or (314) 427-3324 or contact: clones@genomesystems.com or info@genomesystems.com web site: www.genomesystems.com  
Putative full length read

```

FEATURES
Source
vector to vector length is 57.
Location/Qualifiers
1..56
/organism="Glycine max"
/db_xref="taxon:3847"
/clone="GENOME SYSTEMS CLONE ID: Gm-cl037-803"
/clone_id="Gm-cl037"
/tissue_type="fully expanded leaves of greenhouse grown plants"
/dev_stage="2 week old"
/lab_host="DH10B"
/notes="vector: pSPORT1; site_1: NotI; site_2: SalI; This cDNA library was constructed from mRNA isolated from fully expanded leaves of greenhouse grown plants that were 2 weeks old. The library was prepared using the Life Technologies psupertscript cDNA library construction kit. Complementary DNA was synthesized from mRNA using a poly(dT) sequence with a NotI restrictions site. SalI linkers adapters were ligated to the blunt-ended cDNA fragments followed by NotI digestion. The cDNA fragments were directionally cloned into the NotI-SalI restriction site of the pSPORT1 vector. The ligated cDNA fragments were transformed into E.coli Electro-Max DH10B host cells. This library was constructed in the laboratory of Dr. Lila Vodkin by Anu Khanna at the University of Illinois at Urbana-Champaign. email: l-vodkin@uiuc.edu"
15 a 2 c 11 g 28 t
BASE COUNT
ORIGIN

```

Query Match	2.0%	Score 20	DB 10	Length 56
Best Local Similarity	65.9%	Pred. NO. 1.1e+06		
Matches 29	Conservative 0	Mismatches 15	Indels 0	Gaps 0
Qy	846	tgttcagcctctaagatgaaatgctgaactcttcttactt	889	
Db	3	ttttcagctctatagatgatttaattgatttttgatttactt	46	

RESULT	15		
AF149647			
LOCUS	56 bp	DNA	GSS
DEFINITION	AF149647 Human chromosome 18q21 from exon-trapping Homo sapiens genomic clone 5ml0, DNA sequence.		12-JUN-2000
ACCESSION	AF149647		
VERSION	AF149647.1	GI:8485985	
KEYWORDS	GSS.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
AUTHORS	1 (bases 1 to 56)		
TITLE	Chen,H., Huo,Y., Patel,S., Zhu,X., Swift-Scanlan,T., Reeves,R.H., DePaulo,R. Jr., Ross,C.A. and McInnis,M.G.		
JOURNAL	Gene identification using exon amplification on human chromosome 18q21: implications for bipolar disorder		
MEDLINE	Mol. Psychiatry 5 (5), 502-509 (2000)		
COMMENT	20485132		
	Contact: Chen H		
	Psychiatry and Behavioral Sciences		
	Johns Hopkins University School of Medicine		
	600 N. Wolfe Street, Baltimore, MD 21287, USA		
	Email: hcw@chlink.welch.jhu.edu		
	Class: exon-trapped.		

BASE COUNT	24 a	12 c	7 g	13 t
ORIGIN				





alignment\_block:  
 US-09-439-311-2 x AAA86891/rev ..  
 Align seg 1/1 to reverse of: AAA86891 from: 1 to: 30

97 GlnAspGlyGlnSerLeuLysThrArgThr 106  
 ||||||||||||||||||||||||||||  
 30 CACAGTGGACAAAGTTTAAAAACAAGAACT 1

seq\_name: /SID52/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA86892

seq\_documentation\_block:  
 ID AAA86892 standard; DNA; 30 BP.  
 AC AAA86892;  
 XX  
 DT 15-JAN-2001 (first entry)  
 DE Probe to Campylobacter jejuni.  
 XX  
 XX Detection; nucleic acid hybrid; depolymerisation; analysis; SNP;  
 KM single nucleotide polymorphism; identification; viral load; probe;  
 KM genotyping; medical marker diagnostic; primer; target; mutation;  
 KM genetic disease; ss.  
 XX  
 XX Campylobacter jejuni.  
 OS  
 XX MO200049180-A1.  
 PI  
 XX  
 PD 24-AUG-2000.  
 XX  
 XX 18-FEB-2000; 2000WO-US04242.  
 PF  
 XX 18-FEB-1999; 99US-0252436.  
 PR 21-JUL-1999; 99US-0358972.  
 PR 25-AUG-1999; 99US-0383316.  
 XX  
 PA (PROM-) PROMEGA CORP.  
 PI Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
 PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
 XX  
 DR WPI: 2000-565377/52.  
 XX  
 XX Determining presence or absence of a predetermined endogenous nucleic  
 PT acid sequence by using an enzyme that depolymerizes the 3' end of an  
 PT oligonucleotide probe hybridized to a target sequence to release  
 PT identifier nucleotides -  
 XX  
 XX Example: Page 321; 389pp; English.  
 PS  
 XX The present invention describes a method (M1) for determining the  
 CC presence or absence of a predetermined endogenous nucleic acid target  
 CC sequence (ENAT). The method comprises hybridizing a probe having an  
 CC identifier nucleotide (IN) with ENAT which is treated with an enzyme  
 CC that depolymerizes the 3' end of hybridised NA to release the INs.  
 CC M1 is used for determining the number of known sequence repeats present  
 CC in a nucleic acid target sequence in a nucleic acid sample. The method  
 CC is also useful for determining whether a nucleic acid target sequence in  
 CC a sample is an allele from a homozygous or heterozygous locus. The  
 CC method is also useful for detection of mutations, translocations and  
 CC SNPs in nucleic acids (including those associated with genetic disease),  
 CC determination of viral load, species identification, sample  
 CC contamination, and analysis of forensic samples. AAA86791 to AAA87079  
 CC and AAA812817 represent sequence which are used in the exemplification of  
 CC the present invention.  
 CC N.B. There is a discrepancy between the SEQ ID NO: and sequences given  
 CC in the examples, and the SEQ ID NO: and sequences given in the sequence  
 CC listing from the present invention.  
 XX  
 XX Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other:

alignment\_scores:  
 Quality: 50.00 Length: 10  
 Ratio: 5.000 Gaps: 0  
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment\_block:  
 US-09-439-311-2 x AAA86892 ..  
 Align seg 1/1 to: AAA86892 from: 1 to: 30

97 GlnAspGlyGlnSerLeuLysThrArgThr 106  
 ||||||||||||||||||||||||||||  
 1 CACAGTGGACAAAGTTTAAAAACAAGAACT 30

seq\_name: /SID52/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA93188

seq\_documentation\_block:  
 ID AAA93188 standard; DNA; 30 BP.  
 AC AAA93188;  
 XX  
 DT 11-JAN-2001 (first entry)  
 DE Campylobacter jejuni interrogation probe 11451.  
 XX  
 XX Campylobacter jejuni; nucleic acid detection; genomic typing;  
 KM mutation detection; viral load determination; species identification;  
 KM forensic analysis; probe; ss.  
 XX  
 XX Campylobacter jejuni.  
 OS  
 XX MO200049179-A1.  
 PI  
 XX  
 PD 24-AUG-2000.  
 XX  
 XX 18-FEB-2000; 2000WO-US04176.  
 PF  
 XX 18-FEB-1999; 99US-0252436.  
 PR 21-JUL-1999; 99US-0358972.  
 PR 27-SEP-1999; 99US-0406147.  
 XX  
 PA (PROM-) PROMEGA CORP.  
 PI Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
 PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
 XX  
 DR WPI: 2000-549282/50.  
 XX  
 XX Detecting the presence of predetermined exogenous nucleic acid target  
 PT sequence useful for e.g. genotyping, comprises depolymerizing the 3'  
 PT end of an oligonucleotide probe hybridized to a nucleic acid target  
 PT sequence -  
 XX  
 XX Claim 47: Page 187; 230pp; English.  
 PS  
 XX The present sequence is an interrogation probe which was used to detect a  
 CC segment of the genome of Campylobacter jejuni. This was performed as part  
 CC of a method for determining the presence of a known exogenous nucleic  
 CC acid target sequence in a nucleic acid sample. The method comprises  
 CC admixing a treated sample with a depolymerising enzyme which releases one  
 CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
 CC The method is used for assaying nucleic acids for a particular native or  
 CC mutant sequence, and for genomic typing. It is useful for detecting  
 CC mutations, translocations, and single nucleotide polymorphisms,  
 CC determination of viral load, species identification, detection of sample  
 CC contamination, and analysis of forensic samples. Compared with previous  
 CC methods of detecting nucleic acid hybrids, the new method has higher  
 CC sensitivity without the need for radiochemicals or electrophoresis. It is  
 CC quantitative, highly reproducible and can be automated. The method can  
 CC reliably detect as few as 10 copies of a virus in a sample, and is  
 CC capable of providing multiple analyses in a single assay (multiplex  
 CC assay).

XX  
SQ Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;

# alignment\_scores:

Quality: 50.00 Length: 10  
Ratio: 5.000 Gaps: 0  
Percent Similarity: 100.000 Percent Identity: 100.000

# alignment\_block:

US-09-439-311-2 x AAA93188/rev ..

Align seg 1/1 to reverse of: AAA93188 from: 1 to: 30

97 GlnAspGlyGlnSerLeuLysThrArgThr 106  
|||||  
30 CAAGATGACAAAGTTAAAAACAAGAACT 1

seq\_name: /SID22/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA93190

# seq\_documentation\_block:

ID AAA93190 standard; DNA: 30 BP.

XX AAA93190;

XX 11-JAN-2001 (first entry)

XX Campylobacter jejuni Interrogation probe 11450.

XX Campylobacter jejuni; nucleic acid detection; genomic typing;

KW mutation detection; viral load determination; species identification;

XX forensic analysis; probe; ss.

XX Campylobacter jejuni.

XX WO200049179-A1.

XX 24-AUG-2000.

XX 18-FEB-2000; 2000MO-US04176.

XX 18-FEB-1999; 99US-0252436.

XX 21-JUL-1999; 99US-0358972.

XX 27-SEP-1999; 99US-0406147.

XX (PROM-) PROMEGA CORP.

XX Shultz JM, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;

XX Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;

XX WPI; 2000-549282/50.

XX Claim 47; Page 187; 230pp; English.

CC The present sequence is an interrogation probe which was used to detect a  
CC segment of the genome of Campylobacter jejuni. This was performed as part  
CC of a method for determining the presence of a known exogenous nucleic  
CC acid target sequence in a nucleic acid sample. The method comprises  
CC admixing a treated sample with a depolymerising enzyme which releases one  
CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
CC The method is used for assaying nucleic acids for a particular native or  
CC mutant sequence, and for genomic typing. It is useful for detecting  
CC mutations, translocations, and single nucleotide polymorphisms,  
CC determination of viral load, species identification, detection of sample  
CC contamination, and analysis of forensic samples. Compared with previous  
CC methods of detecting nucleic acid hybrids, the new method has higher  
CC sensitivity without the need for radiochemicals or electrophoresis. It is  
CC quantitative, highly reproducible and can be automated. The method can

CC reliably detect as few as 10 copies of a virus in a sample, and is  
CC capable of providing multiple analyses in a single assay (multiplex  
CC assay).

XX  
SQ Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;

# alignment\_scores:

Quality: 50.00 Length: 10  
Ratio: 5.000 Gaps: 0  
Percent Similarity: 100.000 Percent Identity: 100.000

alignement\_block:

US-09-439-311-2 x AAA93190 ..

Align seg 1/1 to: AAA93190 from: 1 to: 30

97 GlnAspGlyGlnSerLeuLysThrArgThr 106  
|||||  
1 CAAGATGACAAAGTTAAAAACAAGAACT 30

seq\_name: /SID22/gcgdata/geneseq/geneseqn/NA1998.DAT:AAV23196

# seq\_documentation\_block:

ID AAV23196 standard; DNA: 59 BP.

XX AAV23196;

XX 28-JUL-1998 (first entry)

XX Lactococcus lactis constitutional promoter Cp29.

KW Lactococcus lactis; constitutional promoter; optimise; spacer;

XX artificial promoter library; gene expression; ds.

XX Lactococcus lactis.

XX Synthetic.

XX Lactococcus lactis.

XX Key

XX Location/Qualifiers

XX Promoter

XX 4..59

XX /\*tag= a

XX /standard\_name= "Constitutional promoter"

XX WO9807846-A1.

XX 26-FEB-1998.

XX 25-AUG-1997; 97MO-DK00342.

XX 23-AUG-1996; 96DK-0000886.

XX (JENSEN) JENSEN P. R.

XX Hammer K, Jensen PR;

XX WPI; 1998-179062/16.

XX Claim 28; Page 52; 89pp; English.

CC This is a Lactococcus lactis constitutional promoter sequence used in the  
CC construction of an artificial promoter library of the invention. The  
CC artificial promoter library for a selected organism or group of organisms  
CC comprise a mixture of double-stranded DNA fragments, the sense strands of  
CC which comprise at least half of two consensus sequences of efficient  
CC promoters from the organism or group of organisms and surrounding or  
CC intermediate nucleotide sequences (spacers) of variable length in which  
CC at least 7 nucleotides are selected randomly, with the proviso that  
CC previously known promoter sequences and promoter sequences isolated from  
CC natural sources are not included. This promoter library can be used in a

```

CC convertases or endoproteases that exhibit testis specificity.
CC Antagonists, including antibodies, are useful for inhibiting or
CC antagonizing the function of ZS1G-11. It is possible that ZS1G-11 and
CC its antagonists will be useful as fertility inducing therapeutics.
CC Sequences AAX34800-21 represent PCR primers for amplifying the ZS1G-11
CC DNA.
XX
XX Sequence 51 BP; 17 A; 5 C; 19 G; 10 T; 0 other;
XX
XX
XX alignment_scores:
XX      Quality: 43.00      Length: 16
XX      Ratio: 3.583      Gaps: 0
XX Percent Similarity: 75.000      Percent Identity: 50.000
XX
XX alignment_block:
XX US-09-439-311-2 x AAX34813 ..
XX
XX Align seg 1/1 to: AAX34813 from: 1 to: 51
XX
XX 253 GYVAIAValIIeGlyLySVaIAsPTySerAspGIyAspGIuAaNgly 268
XX      ||||| ::: ||| |||||:::||| ||:::||||
XX      1 GGTGTAAAGCTTGACACAGAGATTACACAGCAGTCATGACAAAGGCT 48
XX
XX seq_name: /SIDS2/gcdata/geneseq/geneseqn/NA1999.DAT:AAX19519
XX
XX seq_documentation_block:
XX ID AAX19519 standard; DNA: 51 BP.
XX
XX AAX19519;
XX
XX 07-JUN-1999 (first entry)
XX
XX Human lipocalin homologue zlipol PCR primer ZC13,735.
XX
XX DE Human lipocalin: testis; mammary gland; breast tumour; zlipol;
XX KW breast cancer; emphysema; skin disease; reproduction; anti-inflammatory;
XX KW antimicrobial; PCR primer; ss.
XX
XX OS Synthetic.
XX OS Homo sapiens.
XX
XX PN MO9907740-A2.
XX
XX 18-FEB-1999.
XX
XX PF 06-AUG-1998; 98MO-US16425.
XX
XX PR 06-AUG-1997; 97US-0054867.
XX
XX PA (ZYMO ) ZYMOGENETICS INC.
XX
XX Conklin DC;
XX
XX MPI: 1999-167367/14.
XX
XX PT New lipocalin homologue designated zlipol - whose expression is
XX PT restricted to testis and mammary gland tissues, particularly breast
XX PT tumour tissue, used to, e.g. predict tumour aggressiveness.
XX
XX Example 5; Page 89; 94pp; English.
XX
XX
XX The present sequence represents a PCR primer for lipocalin homologue,
XX zlipol. The lipocalin homologue, zlipol, is specifically expressed in
XX testis and mammary gland, particularly breast tumour tissue. Based on
XX this tissue distribution, zlipol may be used as a diagnostic for breast
XX carcinomas and as a tool for predicting tumour aggressiveness. Agonists
XX can be used for transportation of small hydrophobic molecules either in
XX vivo or in vitro, and so are useful in specifically promoting the growth
XX and/or development of testis-specific cell lineages in culture. zlipol
XX can be used to identify inhibitors. zlipol proteins can also be used to
XX prepare antibodies (which can be linked to toxins), and can serve as
XX immunogens. zlipol proteins can be used as a delivery and encapsulation
XX

```



CC system to transport and/or stabilise small lipophilic molecules, e.g. to  
CC protect from gut pH and digestive enzymes. They can also be used to bind  
CC small fatty acids in blood or tissues to modulate their biological  
CC function, e.g. to transport retinoids or steroids to receptors, in  
CC particular as therapy for breast cancer, emphysema and diseases of the  
CC skin. They may also play an important role in reproduction. Other uses  
CC include anti-inflammatory responses, and antimicrobial activities.  
CC Zilipol nucleic acid sequences may be used for gene therapy to increase  
CC or inhibit zilipol activity, to derive probes and primers, to derive  
CC antisense sequences, and to detect genetic abnormalities.  
XX  
SQ Sequence 51 BP; 17 A; 5 C; 19 G; 10 T; 0 other;

Alignment\_scores:  
Quality: 43.00 Length: 16  
Ratio: 3.583 Gaps: 0  
Percent Similarity: 75.000 Percent Identity: 50.000

Alignment\_block:  
US-09-439-311-2 x AAX19519 ..

Align seg 1/1 to: AAX19519 from: 1 to: 51

253 G1YVAV1Val11eG1yLysVAlAsPTySerAspG1yAspG1uAsnG1y 268  
||||| :||| |||||:||||| |||||:|||||  
1 GGTGAACCTTGACACAGACAGATTACACAGACGATGATGACACAGCT 48

seq\_name: /SIDS2/gcgdata/geneseq/geneseqn/NA1998.DAT:AAV00234

seq\_documentation\_block:  
ID AAV00234 standard; DNA: 50 BP.  
XX  
AC AAV00234:  
XX  
DT 08-JUN-1998 (first entry)  
XX  
DE Tick vasoreactive amine binding protein FS-HBPI reverse PCR primer.  
XX  
XX Female-specific vasoreactive amine binding protein 1; FS-HCPI:  
KW histamine; serotonin; assay; antihistamine; anti-inflammatory;  
KW insect bite; snake bite; scorpion bite; dermatitis; vaccine;  
KW transgenic animal; tick; PCR; primer; ss.  
XX  
OS Synthetic.  
OS Rhipicephalus appendiculatus.  
XX  
PN WO9744451-A2.  
XX  
PD 27-NOV-1997.  
XX  
PF 19-MAY-1997; 97WO-GB01372.  
XX  
PR 18-APR-1997; 97GB-0007844.  
PR 18-MAY-1996; 96GB-0010484.  
XX  
PA (OXFO-) OXFORD VACS LTD.  
XX  
PI Nuttall PA, Paesen GC;  
XX  
DR WPI: 1998-018506/02.  
XX  
PT New vasoreactive amine binding proteins and related nucleic acid,  
PT vectors - transformed cells and transgenic animals, used for  
PT assaying or removing histamine and as antihistamine or  
PT anti-inflammatory agents  
XX  
PS Example 3; Page 20; 44pp; English.  
XX  
CC This reverse primer was used with a forward primer (see AAV00233)  
CC to amplify the coding region (see AAV00227) of Rhipicephalus  
CC appendiculatus female-specific histamine binding protein 1  
CC (FS-HBPI) (see AAW37446), a novel vasoreactive amine binding protein

CC (VABP). The primers were designed so that a SacI site was added  
CC upstream of the start codon, while the stop codon was replaced by  
CC a BamHI site, followed by 6 histidine codons and an SpeI site  
CC comprising a T66 stop codon. The PCR product was ligated into  
CC transfect vector pAC1291, generating plasmid pAC129.1-FS1.HIS.  
CC FS-HBPI was expressed as a histidine-tagged protein in Spodoptera  
CC frugiperda Sf21 ovarian cells using a baculovirus expression system.  
CC VABPs can be used to assay or remove histamine, as an antihistamine  
CC or anti-inflammatory agent, and in vaccines.  
XX  
SQ Sequence 50 BP; 11 A; 8 C; 14 G; 17 T; 0 other;

Alignment\_scores:  
Quality: 41.00 Length: 10  
Ratio: 4.100 Gaps: 0  
Percent Similarity: 100.000 Percent Identity: 70.000

Alignment\_block:  
US-09-439-311-2 x AAV00234 ..

Align seg 1/1 to: AAV00234 from: 1 to: 50

262 SerAspG1yAspG1uAsnG1ySerLeu1le 271  
|||||:|||||:|||||:|||||:|||||:|||||  
7 AGTGATGCTGATGATGATGATGATGCCCTTCG 36

seq\_name: /SIDS2/gcgdata/geneseq/geneseqn/NA2000.DAT:AA296924

seq\_documentation\_block:  
ID AA296924 standard; DNA: 59 BP.  
XX  
AC AA296924:  
XX  
DT 14-APR-2000 (first entry)  
XX  
DE S. cerevisiae gene deletion cassette constructing primer YMR290C-S1.  
XX  
KW Antimycotic; mycosis; immunodepression; AIDS; diabetes; fungicide;  
KW mycel; gene deletion; PCR primer; ss.  
XX  
OS Saccharomyces cerevisiae.  
XX  
PN WO9955907-A2.  
XX  
PD 04-NOV-1999.  
XX  
PF 22-APR-1999; 99WO-EP02722.  
XX  
PR 24-APR-1998; 98EP-0401007.  
PR 11-SEP-1998; 98EP-0402254.  
XX  
PA (HMRI ) HOECHST MARION ROUSSEL.  
XX  
PI Diu-Hercend A, Entian K, Koetter P;  
XX  
DR WPI: 2000-105527/09.  
XX  
PT Identifying antimycotic substances useful for drug preparation and  
PT treatment of mycosis -  
XX  
PS Examples; Page 71; 86pp; English.  
XX  
CC The invention provides a method of screening for antimycotic substances  
CC using essential genes from mycelles or a functionally similar mycelle  
CC gene or the corresponding encoded protein as target. The essential gene  
CC useful for screening antimycotic substances is selected from the  
CC following genes: YML114C, YLR186W, YLR215C, YLR222C, YLR243W, YLR272C,  
CC YLR275W, YLR276C, YLR317W, YLR359W, YLR373C, YLR424W, YLR437C, YLR440C,  
CC YML023C, YML049C, YML077W, YML093W, YML127W, YMR033W, YMR131C,  
CC YMR185W, YMR212C, YMR213W, YMR218C, YMR281W, YMR288W, YMR290C, YMR211W,  
CC YMR049C, YMR134W, YDR196C, YDR299W, YDR355C, YDR407C, YDR416W,  
CC YDR449C, YDR472W, YDR499W, YDR141C, YDR324C, YDR325W, YDR398W, YDR246W,

XX The sequences given in AAT00202-25 and AAT00227-57 represent two groups  
CC of ligands to thrombin. These sequences were isolated using the single  
CC stranded DNA molecules given in AAT00201 and AAT00226 which comprise a  
CC 30N and a 60N variable region, respectively. These ligands were  
CC isolated using systematic evolution of ligands by exponential enrichment  
CC (SELEX). The selection was conducted in a buffer solution at 37 deg. C.  
CC After 12 rounds of selection, no additional improvement in binding was  
CC seen. By studying regions of homology between the isolated ligands, a  
CC truncated ligand of 38 nucleotides (see AAC98403-04) was identified which  
CC retains high affinity binding and inhibits clotting. These ligands are  
CC inhibitors of thrombin and are therefore useful in treating thrombin  
CC mediated conditions and in studying the structure and binding of  
CC thrombin.  
XX  
XX  
SQ Sequence 60 BP; 10 A; 11 C; 29 G; 10 T; 0 other;

Quality:	40.00	Length:	19
Ratio:	2.667	Gaps:	0
Percent Similarity:	78.947	Percent Identity:	42.105

Align seg 1/1 to: AAT0025

Align seg 1/1 to: AT00254 from: 1 to: 60

204 ThrSerValGlyThrCylLeuGLyAlaLeuAlaIugLlLleuSArgAs 220  
::: :::::::::::::::::::: :::  
4 ACCGGCGAGGGCGTtAgGgTTtBgAGGCGTTtggCCCACTgtGTtAgGCACGGA 53  
220 malaasp 222

54 CTCGGAT 60

seq\_name: /SIDS2/gcgsdata/geneseq/geneseqn/NA2001.DAT:AAF70806

ID AAF70806 standard; DNA; 60 BP.

AC AAF70806;

DT 20-APR-2001 (first entry)

DE Thrombin high affinity ligand #53.

KM Ligand; basic fibroblast growth factor; bFGF; gene therapy; vascular;

XX

XX

XX

XX

XX

PR 10-JUN-1991; 91US-0714131.

PR 10-FEB-1994; 94US-0195005.

XX

XX

XX

PT Novel nucleic acid ligands to basic fibroblast growth factor that are  
PT useful as inhibitors of basic fibroblast growth factors and 2'-amino  
PT modified RNA ligands, exhibit increased in vivo stability -

```

XX PS Example 19; Column 59-60: 153pp; English.
CC CC The present invention relates to a purified and isolated non-naturally
XX CC occurring DNA ligands to basic fibroblast growth factor (bFGF).
CC CC The ligands are useful as part of gene therapy treatments and
CC CC for diagnosing pathogenesis of vascular diseases including
CC CC intillation and progression of atherosclerosis, acute coronary
CC CC syndromes, vein graft disease and restenosis following coronary
CC CC angioplasty. The ligands have improved stability in vivo.
XX SO Sequence 60 BP: 10 A; 11 C; 29 G; 10 T; 0 other;

alignment_scores:
    Quality: 40.00      Length: 19
    Ratio: 2.667        Gaps: 0
Percent Similarity: 78.947 Percent Identity: 42.105

alignment_block:
US-09-439-311-2 x AAF70806 ..

Align seg 1/L to: AAF70806 from: 1 to: 60

204 ThisServAlGlyThrGlyLeuGlyAlaLeuAlaGlucIleAsnArgAs 220
||||| :|||::| | ||||| ::| :| :|
4 ACCGGCGAGGGCGTAGGGTTGGAAGCGCTTGCCGATGTGTAGCACGCA 53
220 nalaasp 222
|||||
54 CTCGGAT 60

seq_name= /SIDIS2/gcgcdata/geneseq/geneseqn/A42001.DAT:AAI30690
seq_documentation_block:
ID AAI30690 standard; DNA: 31 BP.
XX AC AAI30690:
XX AA130690:
DT 18-OCT-2001 (first entry)
XX DE Human single nucleotide polymorphism (SNP) ATM 2.
XX KM Human: resequence; genotype: disease; forensic; paternity testing;
KM single nucleotide polymorphism; SNP; ss.
XX OS Homo sapiens.
XX FH Key Location/Qualifiers
FH Variation replace(16.G)
FT FT /tag= a
FT FT /standard_name= "single nucleotide polymorphism"
XX NC WO200166800-A2.
XX PD 13-SEP-2001.
XX PE 07-MAR-2001: 2001MO-US07268.
XX PR 07-MAR-2000: 2000US-0187510.
XX PR 22-MAY-2000: 2000US-0206129.
XX PA (WHED ) WHITEHEAD INST BIOMEDICAL RES.
XX PI Cargill M, Ireland JS, Lander ES;
XX MPI; 2001-522952/57.
PT Nucleic acid molecules from the human genome which include polymorphic
PT sites, useful in methods for predicting the presence, absence or
PT severity of a particular phenotype or disorder (e.g. diabetes)
XX associated with a particular genotype -
XX
```

```

PS Claim 1, Page 101, 145pp: English.
CC The invention relates to the identification of nucleic acid molecules
CC (AAI29513-AAI3114) from the human genome which include polymorphic sites
CC which can predispose individuals to disease. Various genes from a number
CC of individuals were resequenced and single nucleotide polymorphisms
CC (SNPs) in these genes discovered. The method is useful for predicting the
CC presence, absence or severity of a particular phenotype or disorder (e.g.
CC diabetes) associated with a particular genotype. The nucleic acids
CC containing the polymorphic sites may be useful in forensics and paternity
CC testing.
XX
SO Sequence 31 BP; 12 A; 4 C; 7 G; 8 T; 0 other;

alignment_scores:
    Quality: 39.00      Length: 9
    Ratio: 4.333      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 77.778

alignment_block:
US-09-439-311-2 x AAI30690 ..

Align seg 1/1 to: AAI30690 from: 1 to: 31

139 ThrAengIngIupheGInIleGlySer 147
|||||:|||||:|||||F||||
5 ACAATGAGCAATTCAGCAATGTGTTCC 31

seq_name: /SIDS2/gcgdata/geneseq/geneseqn/AAI1998.DAT:AAV56429

seq_documentation_block:
ID ID AAV56429 standard; DNA; 47 BP.
XX
AC AAV56429;
XX
DT 20-NOV-1998 (first entry) /
XX
DE Human ICAM-R cDNA primer #27.

Intercellular adhesion molecule; ICAM-R; human; modulator; 14.3.3 family;
HSL-beta; tubulin; inhibitor; stimulator; effector; immune response;
inflammation; disorder; T cell activation; macrophage; Crohn's disease;
adult respiratory distress syndrome; stroke; multiple sclerosis; asthma;
rheumatoid arthritis; tumour growth; human immune deficiency virus;
infection; diabetes; graft vs. host disease; passive immunisation;
primer; ss.
XX
XX Synthetic.
XX OS Homo sapiens.
XX OS
XX
XX US5773218-A.
XX
XX 30-JUN-1998.
XX
XX 07-JUN-1995; 95US-0482882.
XX
XX 05-AUG-1994; 94US-0286754.
XX 27-JAN-1992; 92US-0827689.
XX 26-MAY-1992; 92US-0889724.
XX 05-JUN-1992; 92US-0894061.
XX 22-JAN-1993; 93US-0009266.
XX 26-JAN-1993; 93WO-US00787.
XX 05-AUG-1993; 93US-0102852.
XX 07-JUN-1995; 95US-0482882.
XX
XX (ICOS-) ICOS CORP.
XX
XX Gallatin WM, Vazeux R;
XX
XX WPI: 1998-386989/33.
XX
XX Identifying compounds that modulate interaction of intercellular

```

PT adhesion molecule R - with ligands HSI-beta and tubulin using  
PT two-hybrid assay, useful for treating inflammation, T cell  
PT activation etc.  
XX  
PS Example 13; Column 135-136; 108bp; English.  
XX  
CC AAV56429-V56434 are primers used in the isolation of a novel human  
CC intercellular adhesion molecule, ICAM-R. This sequence is used in a  
CC method which investigates modulators of the interaction between ICAM-R  
CC and the 14.3.3 family member HSI-beta and tubulin. An anti-ICAM-R  
CC antibody optionally coupled to toxin or radionuclide, or an ICAM-R  
CC peptide can block, inhibit or stimulate ligand/receptor interactions  
CC involving ICAM-R, particularly its effector functions involved in  
CC (non)specific immune responses. ICAM-R related agents may be used to  
CC treat or monitor inflammation, disorders involving T cell activation or  
CC macrophages, e.g. adult respiratory distress syndrome, stroke, Crohn's  
CC disease, multiple sclerosis, rheumatoid arthritis, asthma, tumour  
CC growth, human immune deficiency virus infection, diabetes, graft vs. host  
CC disease and many others. Antibodies may also be used for passive  
CC immunisation, for purifying, detecting or quantifying ICAM-R and for  
CC identifying ICAM-R expressing cells.  
XX  
SQ Sequence 47 BP; 9 A; 21 C; 7 G; 10 T; 0 other:  
  
alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667  
  
alignment\_block:  
US-09-439-311-2 x AAV56429/rev ..  
  
Align seg 1/1 to reverse of: AAV56429 from: 1 to: 47  
  
169 ArgPhegluThnGlySerGlnSerPheSerSergly 180  
||||:||||||||| |||:|||||  
44 AGGATGGAGACTGGGTCACGACGATTGGGAGTGA 9  
  
seq\_name: /SIDS2/gcgdata/geneseq/geneseqn/NA1999.DAT:AAV21884  
  
seq\_documentation\_block:  
ID AAV21884 standard; DNA: 47 BP.  
XX  
AC AAV21884:  
XX  
DT 14-MAY-1999 (first entry)  
XX  
DE Primer for antibody against ICAM-R.  
XX  
KW ICAM: immunoglobulin-like loop; intercellular adhesion molecule receptor;  
KW alpha d/CD18; antibody; immunisation; inflammatory response; asthma;  
KW tumour growth; viral infection; therapy; primer; ss.  
XX  
OS Synthetic.  
OS Mus sp.  
XX  
PN US5880268-A.  
XX  
PD 09-MAR-1999.  
XX  
PF 07-JUN-1995; 95US-0483932.  
XX  
OS 05-AUG-1994; 94US-0286754.  
PR 27-JAN-1992; 92US-0827689.  
PR 26-MAY-1992; 92US-0889724.  
PR 05-JUN-1992; 92US-0894061.  
PR 22-JAN-1993; 93US-0009266.  
PR 26-JAN-1993; 93MO-US00787.  
PR 05-AUG-1993; 93US-0102852.  
PR 07-JUN-1995; 95US-0483932.  
XX  
PA (ICOS-) ICOS CORP.

XX  
PI Gallatin WM, Vazeux R;  
XX  
DR WPI; 1999-204041/17.  
XX  
PT New intercellular adhesion molecule receptor (ICAM-R) specific  
PT antibodies - useful for modulating ligand/receptor binding and  
PT biological activities involving ICAM-R, especially those of the  
PT specific and non-specific immune systems  
XX  
PS Example 13; Column 41; 108bp; English.  
XX  
CC This sequence is a primer for DNA encoding an antibody specific for  
CC ICAM-R. The invention relates to antibodies (Ab) which bind specifically  
CC to the intercellular adhesion molecule receptor (ICAM-R), inhibiting the  
CC interaction between ICAM-R and alpha d/CD18. Abs with specific ICAM-R  
CC binding are useful in compositions for immunisation, and for purifying  
CC ICAM-R polypeptides and identifying cells expressing ICAM-R on their cell  
CC surface, modulating ligand/receptor binding and biological activities  
CC involving ICAM-R, especially inflammatory responses of the specific  
CC immune system, the non-specific immune system, monitoring and treating  
CC asthma, tumour growth, and/or metastasis, and viral infection (e.g. HIV  
CC infection). In particular diseases involving an essential T cell  
CC activation (e.g. asthma, psoriasis, diabetes, graft vs. host disease,  
CC tissue transplant rejection, and multiple sclerosis) may be treated with  
CC anti-ICAM-R antibodies. The Abs specifically bind to and identify ICAM-R  
CC and disrupt ICAM-R to cell adhesion molecule, especially alpha d/CD18  
CC binding.  
XX  
SQ Sequence 47 BP; 9 A; 21 C; 7 G; 10 T; 0 other:  
  
alignment\_scores:  
Quality: 39.00 Length: 12  
Ratio: 3.900 Gaps: 0  
Percent Similarity: 83.333 Percent Identity: 66.667  
  
alignment\_block:  
US-09-439-311-2 x AAV21884/rev ..  
  
Align seg 1/1 to reverse of: AAV21884 from: 1 to: 47  
  
169 ArgPhegluThnGlySerGlnSerPheSerSergly 180  
||||:||||||||| |||:|||||  
44 AGGATGGAGACTGGGTCACGACGATTGGGAGTGA 9  
  
seq\_name: /SIDS2/gcgdata/geneseq/geneseqn/NA1999.DAT:AAV69197  
  
seq\_documentation\_block:  
ID AAV69197 standard; DNA: 47 BP.  
XX  
AC AAV69197:  
XX  
DT 17-FEB-1999 (first entry)  
XX  
DE Humanised ICR-1.1 antibody V $\kappa$  region DNA mutating oligo 110.  
XX  
KW Intercellular adhesion molecule polypeptide; ICAM-R; humanised; ICR 1.1;  
KW ICR 8.1; monoclonal antibody; therapeutic; inflammatory; asthma; tumour;  
KW graft-versus-host disease; viral infection; toxin; radionuclide;  
KW neovascularisation site; mutagenic; PCR primer; ss.  
XX  
OS Synthetic.  
OS Mus sp.  
XX  
PN US5837822-A.  
XX  
PD 17-NOV-1998.  
XX  
PF 07-JUN-1995; 95US-0487113.  
PR 07-JUN-1995; 95US-0487113.  
PR 27-JAN-1992; 92US-0827689.  
XX

PR 26-MAY-1992: 92US-0889724.  
 PR 05-JUN-1992: 92US-0894061.  
 PR 22-JAN-1993: 93US-0009266.  
 PR 26-JAN-1993: 93MO-US00787.  
 PR 05-AUG-1993: 93US-0102852.

XX (ICOS-) ICOS CORP.  
 XX

PI Gallatin NM, Vazeux R;  
 XX

DR WPI: 1999-023535/02.  
 XX

XX Humanised antibodies specific for intercellular adhesion molecule  
 PR polypeptide - useful for therapeutic or diagnostic purposes  
 XX

PS Example 13: Column 42: 116pp; English.

CC The invention relates to humanised ICR 1.1 and ICR 8.1 antibodies  
 CC targeted to the human intercellular adhesion molecule polypeptide  
 CC (ICAM-R) polypeptide. Antibodies specific for ICAM-Rs are potentially  
 CC useful as therapeutic compounds, for treating e.g. immune-mediated  
 CC inflammatory conditions (e.g. graft-versus-host disease), asthma,  
 CC tumours or viral infections. Monoclonal antibodies specific for ICAM-R,  
 CC or their conjugates formed with e.g. toxins or radionuclides are useful  
 CC for therapeutically targeting or detecting neovascularisation sites.  
 CC PCR mutagenic oligos AAV69197 and AAV69198 are used in the construction  
 CC of the V<sub>K</sub> region of the humanised antibody ICR-1.1.  
 XX

SQ Sequence 47 BP: 9 A; 21 C; 7 G; 10 T; 0 other:

# alignment\_scores:

Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

# alignment\_block:

US-09-439-311-2 x AAV69197/rev ..

Align seg 1/1 to reverse of: AAV69197 from: 1 to: 47

169 ArgPheGluThrGlySerGlnSerPheSerSercly 180  
 |||:::||||||| |||:::|||||  
 44 AGGATGGAGACTGGCTCAGCAGCATTTGGAGTGA 9



OM of: US-09-439-311-2 to: EST:\* out\_format : pfs

Date: Apr 17, 2002 2:43 AM

About: Results were produced by the GenCore software, version 4.5,  
Copyright (c) 1993-2000 Compugen Ltd.

#### Command line parameters:

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-MODEL-frame+ .p2n.model -DEV-rlp
-O/-CGN2.1/USPTO.spool/US09439311/runat_16042002_134010_11677/app_query.fasta_1.395
-DB-EST -OFMT=fastap -SUFFIX=est -GAPOP=12.000 -GAPEXT=4.000
-MINMAPCH=0.100 -LOOPEL=0.000 -LOOPEXT=0.000 -OGAPOP=4.500
-OGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
-FCAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DEL0P=6.000
-DELXT=7.000 -START=1 -MATRIX=blomsum62 -TRANS=human40.cdi
-LIST=45 -DOCALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0
-ALIGN=15 -MODE=LOCAL -OUTFMT=ps -NORM=ext -MINLEN=0 -MAXLEN=60
-USER=US09439311@CGN1_1.3691 -NCPU=6 -ICPU=3 -LONGLOG -NO_XLPHY
-WAIT -THREADS=1
```

#### Search information block:

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Query: US-09-439-311-2
Query length: 333
Database: EST:*
Database sequences: 11351937
Database length: 1077921985
Search time (sec): 1467.110000
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#### score\_list:

Sequence	Strd	Orig	ZScore	Escore	Len	Documentation
gb_estl:AU104260	+	48.00	103.91	1.3e+04	50	AU104260 AU104260 Sugano Homo sa
gb_gss:B04096	+	45.00	96.72	1.2e+04	60	B04096 CSRL-25g1-u CSRL flow sor
gb_gss:AZ669793	+	33.00	88.62	9.0e+04	44	AZ669793 IM0283E04R Mouse 10kb F
gb_estl:AU107968	+	33.00	87.43	1.0e+05	50	AU107968 AU107968 Sugano Homo sa
gb_estl:W20078	+	33.00	87.07	1.1e+05	52	W20078 zb40e03.r1 Soares_parity
gb_gss:AZ2921603	+	33.00	86.21	1.2e+05	57	AZ2921603 1006030F10.x1 1006 - R
gb_gss:AZ298589	+	38.50	85.80	1.3e+05	54	AZ298589 2M0285D08R Mouse 10kb F
gb_estl:AU106648	+	38.00	85.60	1.3e+05	50	AU106648 AU106648 Sugano Homo sa
gb_estl:AU106648	+	38.00	84.55	1.3e+05	56	AU106648 AU106648 Sugano Homo sa
gb_gss:AZ654882	+	38.00	84.38	1.5e+05	57	AZ654882 IM0529N22F Human adult
gb_estl:AA628048	+	37.00	83.90	1.6e+05	33	AA628048 ng62f05.s1 NCI_CGAP_OV4
gb_gss:AZ429658	+	37.00	83.92	1.0e+05	60	AZ429658 IM0213A18R Mouse 10kb F
gb_estl:AU1595782	+	37.00	83.96	1.6e+05	49	AU1595782 uj69c12.x1 Sugano mouse
gb_estl:AU102738	+	37.00	83.77	1.7e+05	50	AU102738 AU102738 Sugano Homo sa
gb_gss:CN504000	+	37.00	83.77	1.7e+05	50	AU102738 AU102738 Sugano Homo sa
gb_estl:AA664752	+	37.00	82.55	2.0e+05	57	AA664752 tetraodon nigroviridis
gb_estl:AU1320166	+	37.00	82.47	2.3e+05	58	AU1320166 c1e11m.r1 Neurospora c
gb_gss:AZ2996603	+	36.50	81.47	2.3e+05	58	AZ2996603 2M0282P18R Mouse 10kb F
gb_gss:AZ615427	+	36.00	83.78	1.7e+05	41	AZ615427 IM0444G18R Mouse 10kb F
gb_estl:AU1522735	+	36.00	82.71	1.9e+05	46	AU1522735 fb61e01.x1 Zebrafish wa
gb_gss:AZ228471	+	36.00	82.32	2.0e+05	48	AZ228471 IM0052F16F Mouse 10kb F
gb_estl:AU105900	+	36.00	81.94	2.1e+05	50	AU105900 AU105900 Sugano Homo sa
gb_estl:AU1905591	+	36.00	81.75	2.2e+05	51	AU1905591 CM-BR094-050299-117 BRG
gb_gss:AZ402345	+	36.00	81.75	2.2e+05	52	AZ402345 IM0169B10R Mouse 10kb F
gb_estl:BB043282	+	36.00	81.05	2.4e+05	55	BB043282 h448h12.y1 NCI_CGAP_OV3
gb_estl:AA673303	+	36.00	81.05	2.4e+05	55	AA673303 vm69f08.r1 Barstead mol
gb_estl:AA711965	+	36.00	81.05	2.4e+05	55	AA711965 vu29c05.x1 Zebrafish mou
gb_estl:AU1444123	+	36.00	81.05	2.4e+05	55	AU1444123 fb26b06.x1 Zebrafish wa
gb_estl:AU1831322	+	36.00	81.05	2.4e+05	55	AU1831322 w182d10.x1 NCI_CGAP_LY
gb_gss:AZ212625	+	35.50	79.48	2.9e+05	59	AZ212625 IM0168L07F Mouse 10kb F
gb_estl:AU1633469	+	35.00	81.51	2.2e+05	43	AU1633469 th62d06.x1 NCI_CGAP_OV2
gb_estl:W53880	+	35.00	80.29	2.2e+05	43	W53880 md03h05.r1 Soares mouse e
gb_estl:AU007710	+	35.00	80.29	2.6e+05	49	AU007710 AU007710 Schizosaccharc
gb_estl:AA400193	+	35.00	80.11	2.7e+05	50	AA400193 zu64e08.s1 Soares_testi
gb_estl:AU106926	+	35.00	79.74	2.8e+05	52	AU106926 AU106926 Sugano Homo sa
gb_estl:AU1653077	+	35.00	79.74	2.8e+05	52	AU1653077 tw33c01.x1 NCI_CGAP_G6
gb_estl:AA449400	+	35.00	79.74	2.8e+05	52	AA449400 z36h11.x1 NCI_CGAP_UC1
gb_estl:BB654043	+	35.00	78.89	3.1e+05	57	BB654043 sad50f03.y2 Gm-cl075 GI
gb_estl:BG099588	+	35.00	78.89	3.1e+05	57	BG099588 nag54e01.x1 NCI_CGAP_Cd

seq\_name: gb\_estl:AU104260

seq\_documentation\_block:

LOCUS AU104260 50 bp mRNA EST 05-APR-2001

DEFINITION AU104260 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone

ACCESSION HEPI15388 mRNA sequence.

VERSION AU104260

KEYWORDS AU104260.1 GI:13553781

SOURCE EST.

ORGANISM human.

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 50)

AUTHORS Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata

,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,T., Morishita,S., Okubo

,K., Suyama,A. and Sugano,S.

Fine structural analysis of transcription start sites of human

mRNAs using full-length enriched and 5'-end enriched cDNA libraries

Unpublished (2001)

CONTACT Yutaka Suzuki

DEPARTMENT Department of Virology

INSTITUTE Institute of Medical Science, University of Tokyo

4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan

EMAIL: yusuzuki@ims.u-tokyo.ac.jp

SUZUKI,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano

S. Construction and characterization of a full length-enriched and

a 5'-end enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES

source

1..50

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="HEPI15388"

/clone\_ltb="Sugano Homo sapiens cDNA library"

BASE COUNT 17 a 12 c 11 g 10 t

ORIGIN

alignment\_scores:

Quality: 48.00 Length: 14

Ratio: 4.364 Gaps: 0

Percent Similarity: 78.571 Percent Identity: 71.429

alignment\_block:

US-09-439-311-2 x AU104260 ..

Align seg 1/1 to: AU104260 from: 1 to: 50

143 PheGlnIleGlySerSerAsnGlnThrIleTysAlaSer 156

|||||:|||||

4 TTTCAGCTTGACACTGTTCCAAATCAGACCAACGACAGC 45

seq\_name: gb\_gss:B04096

seq\_documentation\_block:

LOCUS B04096 60 bp DNA GSS 13-JUL-1996

DEFINITION CSRL-25g1-u CSRL flow sorted chromosome 11 specific cosmid Homo

sapiens genomic clone CSRL-25g1, DNA sequence.

ACCESSION B04096

VERSION B04096.1 GI:1413374

KEYWORDS GSS.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 60)

EVANS,G.A., Burbee,D., Davies,C., Hahner,L., Oliver,T., Gilbert,M.,

Jones,D., Ward,T., Gillilan,E., Schlegmann,J., Probst,S., Harris

TITLE  
JOURNAL  
COMMENT

J., DeFord, J., McFarland, J., Burzinski, K., Khan, M., Kupfer, K. and Garner, H.R.  
Genomic Sequence Sampled Map of Chromosome 11  
Unpublished (1996)  
Contact: Evans GA, Shane Probst  
Modermott Center for Human Growth and Development  
University of Texas Southwestern Medical Center At Dallas  
5323 Harry Hines Blvd, Dallas TX 75235-8591  
Tel: 214-648-1600  
Fax: 214-648-1666  
Email: gevarns@utsw.swmed.edu, shane@modermott.swmed.edu

FEATURES  
source  
Class: cosmid ends  
High quality sequence stop: 60.  
Location/Qualifiers  
1. 60  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="CSRL-25g1"  
/clone\_lib="CSRL flow sorted Chromosome 11 specific cosmid"  
/sex="female"  
/cell\_type="chimeric hamster somatic cell hybrid"  
/note="Vector: sCos-1, Human Chromosome 11 specific cosmid library prepared from flow sorted human Chromosome 11 derived from Chinese Hamster Ovary (CHO) monochromosomal somatic cell hybrid, J1"

BASE COUNT 16 a 16 c 5 g 22 t 1 others  
ORIGIN

alignment\_scores:  
Quality: 45.00 Length: 17  
Ratio: 3.000 Gaps: 0  
Percent Similarity: 88.235 Percent Identity: 52.941

alignment\_block:  
US-09-439-311-2 x B04096 ..  
Align seg 1/1 to: B04096 from: 1 to: 60

112 TleasnaTleuMetGluGluLeuAspAsnIleAlaAsnThrSerp 128  
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
4 ATTACGCTCTCATTTGAGACCTCCCTTCATTTCNAATACAAATAGTTT 53

128 e 128  
1  
54 C 54

seq\_name: gb\_gss:A2469793

seq\_documentation\_block:  
LOCUS A2469793 44 bp DNA GSS 04-OCT-2000  
DEFINITION IM0283F04R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
clone UUGC1M0283F04 R. DNA sequence.

ACCESSION A2469793  
VERSION A2469793.1 GI:10627918  
KEYWORDS GSS.  
SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 44)  
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A. and Wright, D., Weis, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts  
JOURNAL Unpublished (2000)  
COMMENT Contact: Robert B. Weis  
University of Utah Genome Center  
University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SIC, UT 84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00  
Plate: 0283 row: F column: 04  
Seq primer: CACCAAGGAAACAGCTATGACC  
Class: plasmid ends  
High quality sequence stop: 44.  
Location/Qualifiers  
1. 44  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone\_lib="UUGC1M0283F04"  
/clone\_lib="Mouse 10kb plasmid UUGC1M library"  
/sex="Male"  
/lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
/note="Vector: pMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g114732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 9 a 12 c 12 g 11 t  
ORIGIN

alignment\_scores:  
Quality: 39.00 Length: 13  
Ratio: 3.545 Gaps: 0  
Percent Similarity: 84.615 Percent Identity: 53.846

alignment\_block:  
US-09-439-311-2 x A2469793 ..  
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5 GGGGCTGGCATTGCTCTCATATGACAACTTTGGCAAGCTC 43

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seq\_documentation\_block:  
LOCUS A0107968 50 bp mRNA EST 05-APR-2001  
DEFINITION A0107968 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone  
KRT11.118, mRNA sequence.

ACCESSION A0107968  
VERSION A0107968.1 GI:13557490  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 50)  
Suzuki, Y., Tsunoda, T., Taira, H., Mizushima-Sugano, J., Sese, J., Hata, H., Ota, T., Isogai, T., Tanaka, T., Nakamura, Y., Morishita, S., Okubo, K., Suyama, A. and Sugano, S.



```

/clone="IMAGE:306076"
/clone_lib="Scares_Parathyroid_tumor_NbHNA"
/rclone_type="Parathyroid tumor"
/dev_stage="adult"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: parathyroid gland; Vector: pT7T3D (Pharmacia) with a modified polylinker; Site.1: Not I; Site.2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer
[5'-TGTTACCAATCTGAGTGGAGCGCCGACCAATTTTTTTTTTTTTTTTTTTT
TTTTT-3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT7T3D vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by Benito Soares and M. Fatima Bonaldi. RNA from sporadic parathyroid adenomas was kindly provided by Dr. Stephen Marx, National Institute of Diabetes and Digestive and Kidney Diseases, NIH."
BASE COUNT      8 a      7 c      19 g      14 t      4 others
ORIGIN
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    Quality:      39.00      Length:      17
    Ratio:        3.545      *      Gaps:      0
    Percent Similarity: 64.706      Percent Identity: 47.059
alignment_block:
US-09-439-311-2 x W20078      ..
Align seg 1/1 to: W20078 from: 1 to: 52
299 AspGIyArGlyTleYsIleThrgYserIleGlyValGIyAlaGlyI 315
:::|||||:::|||||:::|||||:::|||||:::|||||
2 AATGGTGNAGGTCCTGCACAGATGAGCGCMTGGCTGCTGGTCTGGAT 51
315 e 315
52 T 52
seq_name: gb_gss:A2921603
seq_documentation_block:
LOCUS      A2921603      57 bp      DNA      GSS      20-MAR-2001
DEFINITION      1006030F10.x1 1006 - Rescuem Grid G zea mays genomic, DNA
sequence.
ACCESSION      A2921603
VERSION      A2921603.1 GI:13393406
KEYWORDS      GSS.
SOURCE      Zea mays.
ORGANISM      Zea mays
Eukaryote; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
Clade: Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 57)
Walbot.V.
Maize genomic sequences found using engineered Rescuem transposon
unpublished (2001)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Sequence was trimmed at possible ligation site. Post-ligation
sequence submitted separately.
Plate: 1006030 row: F column: 10
Class: transposon-tagged.
Location/Qualifiers
1..57
/organism="Zea mays"

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/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/clone_lib="1006 - Rescuemu Grid G"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/notes="Organ: leaf; Vector: Rescuemu (engineered from
pBluescript backbone); Site:1: BamHI; Site:2: BglII;
Rescuemu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on Rescuemu, go to the web
site 'www.zmbl.ias.tate.edu' and follow the links for
'Rescuemu.' Grid G was grown at Stanford in 2000. DNA was
extracted from leaf punches, double digested using BamHI
and BglII, and ligated to form circular plasmids. DH10B
cells were transformed and then screened on LB plates with
ampicillin."

BASE COUNT      13 a      18 c      7 g      19 t
ORIGIN

alignment_scores:
    Quality:      39.00      Length:      11
    Ratio:         4.333      Gaps:         0
    Percent Similarity: 81.818      Percent Identity: 54.545

alignment_block:
US-09-439-311-2 x A2921603/rev ..

Align seg 1/1 to reverse of: A2921603 from: 1 to: 57

    309 IIEGlyVAIGlyAlGlylleuH1strGlu 319
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    54 CTGGGGATGGAAGTGGATATCTGCACAGAG 22

seq_name: gb_gss:A2998589

seq_documentation_block:
LOCUS      A2998589      54 bp      DNA      GSS      27-APR-2001
DEFINITION      2M0285D08R Mouse 10kb plasmid UUGC2M library Mus musculus genomic
clone UUGC2M0285D08 R, DNA sequence.
ACCESSION      A2998589
VERSION      A2998589.1 GI:13869816
KEYWORDS      GSS.
SOURCE      house mouse.
ORGANISM      Mus musculus
                Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;
                1 (bases 1 to 54)
REFERENCE      Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
AUTHORS      Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
                ,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
                and Wright,D., Weiss,R.
TITLE      Mouse whole genome scaffolding with paired end reads from 10kb
                plasmid inserts
JOURNAL      Unpublished (2000)
COMMENT      Contact: Robert B. Weiss
                University of Utah Genome Center
                University of Utah
                Rm. 308, Biomedical polymers Research Bldg., 20 S. 2030 E., SLIC, UT
                84112, USA
                Tel: 801 585 5606
                Fax: 801 585 7177
                Email: ddunn@genetics.utah.edu
                Insert Length: 10000 Std Error: 0.00
                Plate: 0285 row: D column: 08
                Seq Primer: CACACAGGAACACCTATGACC
                Class: plasmid ends
                High quality sequence stop: 54.
                location/Qualifiers
                1..54
                /organism="Mus musculus"

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/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_lib="UUGC2M0285D08"
/clone_lib="Mouse 10kb plasmid UUGC2M library"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/notes="Vector: PMD42uv; Purified genomic DNA from M.
musculus C57BL/6J (female) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g114732141gblAF129072.1), a copy-number
inducible derivative of plasmid RL. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT      11 a      2 c      21 g      20 t
ORIGIN

alignment_scores:
    Quality:      38.50      Length:      18
    Ratio:         2.750      Gaps:         1
    Percent Similarity: 77.778      Percent Identity: 50.000

alignment_block:
US-09-439-311-2 x A2998589 ..

Align seg 1/1 to: A2998589 from: 1 to: 54

    194 AApphelyPheapsPserValIleserTnserValGlyThrGlye 210
    ::::::::::::::::::::|||
    10 GATTATTAATTGCAATCTGTGATTT.....GTGGGGTAGGGGCT 50

    210 uGly 211
    :|||
    51 GCGG 54

seq_name: gb_est1:AUI06648

seq_documentation_block:
LOCUS      AUI06648      50 bp      mRNA      EST      05-APR-2001
DEFINITION      AUI06648 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
KAT05523, mRNA sequence.
ACCESSION      AUI06648
VERSION      AUI06648.1 GI:13556169
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
                Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
                1 (bases 1 to 50)
REFERENCE      Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata
AUTHORS      ,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo
                ,K., Suyama,A. and Sugano,S.
TITLE      Fine Structural analysis of transcription start sites of human
                mRNAs using full-length enriched and 5'-end enriched cDNA libraries
JOURNAL      Unpublished (2001)
COMMENT      Contact: Yutaka Suzuki
                Department of Virology
                Institute of Medical Science, University of Tokyo
                4-6-1, Shirokane-dai, Minatoku, Tokyo 108-8639, Japan
                Email: yusuzuki@ims.u-tokyo.ac.jp
                Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano

```

, S. Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

# FEATURES

Location/Qualifiers  
1..50  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_1lb="KAT05523"  
/clone\_1lb="Sugano Homo sapiens cDNA library"

BASE COUNT 10 a 20 c 8 g 12 t  
ORIGIN

## alignment\_scores:

Quality: 38.00 Length: 16  
Ratio: 3.455 Gaps: 0  
Percent Similarity: 68.750 Percent Identity: 56.250

## alignment\_block:

US-09-439-311-2 x AU106648/rev ..

Align seg 1/1 to reverse of: AU106648 from: 1 to: 50

247 Glnaaphaalaileasnglyvallelglylvalspyrser 262

49 CAGGCTGCAGCTGTGTGAGTCAAGGGGAGTACTATCG 2

seq\_name: gb\_est2:C20861

seq\_documentation\_block:

LOCUS C20861 56 bp mRNA 23-OCT-1996  
DEFINITION HUMGS0004926 Human adult (K.Okubo) Homo sapiens cDNA 3', mRNA  
sequence.  
ACCESSION C20861  
VERSION C20861.1 GI:1621971  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 56)  
Okubo, K.  
Bodymap: human gene expression database  
Unpublished (1995)  
Contact: Okubo, K.  
Institute for Molecular and Cellular Biol  
Osaka University  
1-3, Yamada-oka, Suita, Osaka Pref. 565, Japan  
Tel: 06-877-5111(ex.3315)  
Email: kousaku@imcb.osaka-u.ac.jp  
Human Gene Signature, 3'-directed cDNA sequence. We are not  
submitting the same cDNA sequence redundantly to DBJ since 1993.  
For the abundance information of clones with this sequence in this  
library and as well as in other 3'-directed libraries, see  
http://www.imcb.osaka-u.ac.jp/bodymap/. The sequences of the clones  
represented by this GS sequences is also found there.

## FEATURES

source

1..56  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_1lb="Human adult (K.Okubo)"  
/dev\_stage="adult"  
BASE COUNT 19 a 9 c 7 g 19 t 2 others  
ORIGIN

## alignment\_scores:

Quality: 38.00 Length: 18  
Ratio: 2.923 Gaps: 0  
Percent Similarity: 72.222 Percent Identity: 44.444

## alignment\_block:

US-09-439-311-2 x C20861/rev ..

Align seg 1/1 to reverse of: C20861 from: 1 to: 56

215 Gluclulleasnaarganalaasplythrlyleargalarnpheas 231

54 GAACAGATATATCTTCTAATACAGTTGTCTAGATTAAGATTCCNNTTGA 5

231 pval 232

4 GATC 1

seq\_name: gb\_gss:A2654882

seq\_documentation\_block:

LOCUS A2654882 57 bp DNA 14-DEC-2000  
DEFINITION 1M0529N22F Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
clone UUGC1M0529N22 F, DNA sequence.  
ACCESSION A2654882  
VERSION A2654882.1 GI:11792028  
KEYWORDS GSS.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 57)  
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T., Reilly,  
M., Rose, M., Rose, R., Stokes, R., Tiney, A., von Niederhausen, A.  
and Wright, D., Weiss, R.,  
Mouse whole genome scaffolding with paired end reads from 10kb  
plasmid inserts  
Unpublished (2000)  
Contact: Robert B. Weiss  
University of Utah Genome Center  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00  
Plate: 0529 row: N column: 22  
Seq primer: CGTTGTAAACGACGCCAGT  
Class: plasmid ends  
High quality sequence stop: 57.

## FEATURES

source

1..57  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUGC1M0529N22"  
/clone\_1lb="Mouse 10kb plasmid UUGC1M library"  
/sex="Male"  
/lab\_host="E. coli strain XL10-Gold, T1-resistant, F-"  
/note="Vector: PMD2ny; Purified genomic DNA from M.  
musculus C57BL/6J (male) was obtained from the Jackson  
Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA  
was hydrodynamically sheared by repeated passage through a  
0.005 inch orifice at constant velocity. The sheared DNA  
was blunt end-repaired with T4 DNA polymerase and T4  
polynucleotide kinase. Adaptor oligonucleotides were  
ligated to the blunt ends in high molar excess. The  
adapted DNA was purified and size-selected for a 9.5 to  
10.5 kb range using preparative agarose gel  
electrophoresis. Vector DNA was prepared from a derivative  
of PMD42 (g114732114[gb|AF129072.1]), a copy-number  
inducible derivative of plasmid R1. The vector was ligated  
with adaptors complementary to the insert adaptors and  
purified. The sheared, adapted mouse DNA was annealed to  
adapted vector DNA, and transformed into  
chemically-competent E. coli XL10-Gold (Stratagene) cells  
and selected for ampicillin resistance.

BASE COUNT 16 a 23 c 6 g 12 t

## ORIGIN

## alignment\_scores:

Quality: 38.00 Length: 14  
Ratio: 3.800 Gaps: 0  
Percent Similarity: 71.429 Percent Identity: 42.857

## alignment\_block:

US-09-439-311-2 x AZ654882/rev ..

Align seg 1/1 to reverse of: AZ654882 from: 1 to: 57

302 GlyIleLysIleThrGlySerIleGlyAlaGlyAlaGlyIle 315

||||| ||| ::|||::||| |||::|  
43 GGCACTAGCATATTGCTGTGGCATAGCCAAAGCATG 2

seq\_name: gb\_est1:AA628048

seq\_documentation\_block:

LOCUS AA628048 60 bp mRNA EST 31-OCT-1997  
DEFINITION ng62805.s1 NCI-CGAP\_Ov6 Homo sapiens cDNA clone IMAGE:1154625

(HUMAN); mRNA sequence.  
Similar to gb:U07857.14 KD PROTEIN OF SIGNAL RECOGNITION PARTICLE

ACCESSION AA628048  
VERSION AA628048.1 GI:2540047

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 60)  
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgaps-remail.nih.gov

Tissue Procurement: Andrew Berchuck, M.D., Elise Kohn, M.D.,  
Rodrigo F. Chuquib, M.D., Michael R. Emmert-Buck, M.D., Ph.D.

CDNA Library Preparation: David B. Krizman, Ph.D.  
CDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/dbp/Image/Image.html

Trace considered overall poor quality  
Insert Length: 300 Std Error: 0.00

Seq primer: -40m13 fwd. ET from Amersham  
High quality sequence stop: 1.

## FEATURES

source

1..60

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone\_image="1154625"

/clone\_lib="NCI-CGAP\_Ov6"

/sex="female"

/tissue\_type="normal cortical stroma"

/lab\_host="DH10B"

/note="Organ: ovary; Vector: PAMF10; mRNA made from normal  
ovarian cortical stroma, cDNA made by oligo-dT priming.  
Non-directionally cloned. Size-selected on agarose gel,  
average insert size 600 bp."

BASE COUNT 18 a 14 c 12 g 16 t

ORIGIN

## alignment\_scores:

Quality: 38.00 Length: 12  
Ratio: 3.455 Gaps: 0  
Percent Similarity: 91.667 Percent Identity: 58.333

## alignment\_block:

US-09-439-311-2 x AA628048 ..

Align seg 1/1 to: AA628048 from: 1 to: 60

180 GlyThrValGlyLeuThrIleLysAsnTyrAsnGly 191

||||| ::|||::||| |||::|  
7 GGCACTAGCATATTGCTGTGGCATAGCCAAAGCATG 42

seq\_name: gb\_gss:AZ429658

seq\_documentation\_block:

LOCUS AZ429658 33 bp DNA GSS 03-OCT-2000  
DEFINITION 1M0213A18R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
clone UUGC1M0213A18 R, DNA sequence.

ACCESSION AZ429658  
VERSION AZ429658.1 GI:10553671

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 33)  
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Kelly,  
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.  
and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb  
plasmid inserts

Unpublished (2000)

COMMENT Contact: Robert B. Weiss  
University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT  
84112, USA

Tel: 801 585 5606  
Fax: 801 585 7177

Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00

Plate: 0213 row: A column: 18  
Seq primer: CACCAAGGAAACAGCATATGACC

Class: plasmid ends  
High quality sequence stop: 33.

Location/Qualifiers

1..33  
/organism="Mus musculus"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone\_image="UUGC1M0213A18"

/clone\_lib="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab\_host="E. coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42nv. Purified genomic DNA from M.  
musculus C57BL/6J (male) was obtained from the Jackson  
Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA  
was hydrodynamically sheared by repeated passage through a  
0.005 inch orifice at constant velocity. The sheared DNA  
was blunt end-repaired with T4 DNA polymerase and T4  
polynucleotide kinase. Adaptor oligonucleotides were  
ligated to the blunt ends in high molar excess. The DNA  
was ligated to the blunt ends in high molar excess. The  
adapted DNA was purified and size-selected for a 9.5 to  
10.5 kb range using preparative agarose gel  
electrophoresis. Vector DNA was prepared from a derivative  
of pMD42 (g1473211419b/AP12072.1), a copy-number  
inducible derivative of plasmid R1. The vector was ligated  
with adaptors complementary to the insert adaptors and  
purified. The sheared, adapted mouse DNA was annealed to  
chemically-competent E. coli XL10-Gold (Stratagene) cells  
and selected for ampicillin resistance."

BASE COUNT 9 a 5 c 11 g 8 t

ORIGIN

```

Institute of Medical Science). Custom primers for
sequencing 5' end primer CTTCTGGCTCTAAAGCTCG and 3' end
primer CGACCTGCAGCTGACGACA."

BASE COUNT      10 a      12 c      11 g      16 t
ORIGIN

alignment_scores:
    Quality:      37.00      Length:      12
    Ratio:        3.700      Gaps:      0
    Percent Similarity: 83.333      Percent Identity: 58.333

alignment_block:
US-09-439-311-2 x AI595782 ..

Align seg 1/1 to: AI595782 from: 1 to: 49

126 ThSerpheasnnglyLysGlnleuSerglyly 137
||| |||:|||||:||||| ||| ||| |||
7 ACACCTTTTCTGTCGAATGCTAGCTTCTGTGGCG 42

seq_name: gb_est1:AU102738

seq_documentation_block:
LOCUS      AU102738      50 bp      mRNA      EST      05-APR-2001
DEFINITION AU102738 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
HRC13040. mRNA sequence.
ACCESSION  AU102738
VERSION     AU102738.1 GI:13552259
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE  1 (bases 1 to 50)
AUTHORS   Suzuki,Y., Tsunoda,T., Taira,H., Mitsuhashi-Sugano,J., Sese,J., Hatake
            ,H., Ota,T., Iisaga,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo
            ,K., Suyama,A. and Sugano,S.
TITLE      Fine structural analysis of transcription start sites of human
            mRNAs using full-length enriched and 5'-end enriched cDNA libraries
            Unpublished (2001)
JOURNAL    Contact: Yutaka Suzuki
            Department of Virology
            Institute of Medical Science, University of Tokyo
            4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
            Email: yusuzuki@ems.u-tokyo.ac.jp
            Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
            ,S. Construction and characterization of a full length-enriched and
            a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES
     source             1..50
                        /organism="Homo sapiens"
                        /db_xref="taxon:9606"
                        /clone="HRC13040"
BASE COUNT      7 a      16 c      18 g      9 t
ORIGIN

alignment_scores:
    Quality:      37.00      Length:      14
    Ratio:        2.846      Gaps:      0
    Percent Similarity: 92.857      Percent Identity: 50.000

alignment_block:
US-09-439-311-2 x AU102738/rev ..

Align seg 1/1 to reverse of: AU102738 from: 1 to: 50

93 ThrGlnAlaIaGlnAspGlyInserLeuLysThrArgThr 106
||||:|||||:||||| |||:|||||:||||| |||
42 ACTAGGGCGCGCGTACAGGAGCTCATTCTCTCCGACG 1


```

```

seq_name: gb_est1:AU102739
seq_documentation_block:
LOCUS      AU102739          50 bp      mRNA          EST          05-APR-2001
DEFINITION AU102739 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
            HRC13119, mRNA sequence.
ACCESSION  AU102739
VERSION    AU102739.1  GI:13552260
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 50)
AUTHORS   Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata
            ,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo
            ,K., Suyama,A. and Sugano,S.
            Fine structural analysis of transcription start sites of human
            mRNAs using full-length enriched and 5'-end enriched cDNA libraries
            Unpublished (2001)
JOURNAL    Contact: Yutaka Suzuki
            Department of Virology
            Institute of Medical Science, University of Tokyo
            4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
            Email: yusuzuki@ims.u-tokyo.ac.jp
            Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
            ,S. Construction and characterization of a full length-enriched and
            a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES   location/Qualifiers
            source
            1..50
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /clone_lib="HRC13119"
            /clone_lib="Sugano Homo sapiens cDNA library"
BASE COUNT 7 a 16 c 18 g 9 t
ORIGIN
alignment_scores:
            Quality: 37.00      Length: 14
            Ratio: 2.846      Gaps: 0
            Percent Similarity: 92.857      Percent Identity: 50.000
alignment_block:
US-09-439-311-2 x AU102739/rev ..
Align seg 1/1 to reverse of: AU102739 from: 1 to: 50
93 ThrglnAlaAlaGlnAspGlyGlnSerLeuLysThrArgThr 106
|||||
42 ACTAGGCCGCGCGCTACAGGAGCTCATTTCTCCCGCAGC 1

```